

GenCore version 5.1.4.p5.4578  
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OM protein - protein search, using sw model

Run on: May 13, 2003, 14:00:14 : Search time 35 Seconds  
(without alignments)  
167.515 Million cell updates/sec

Title: SEQ1-EDITED  
Perfect score: 199  
Sequence: 1 ANSFLXLRHSLXRCIX.....XXAKXIFZVDVDTLAFWSKH 44

Scoring table: BIOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

A\_Geneseq\_101002:\*

- 1: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1980.DAT:\*
- 2: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1981.DAT:\*
- 3: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1982.DAT:\*
- 4: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1983.DAT:\*
- 5: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1984.DAT:\*
- 6: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1985.DAT:\*
- 7: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1986.DAT:\*
- 8: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1987.DAT:\*
- 9: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1988.DAT:\*
- 10: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1989.DAT:\*
- 11: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1990.DAT:\*
- 12: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1991.DAT:\*
- 13: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1992.DAT:\*
- 14: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1993.DAT:\*
- 15: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1994.DAT:\*
- 16: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1995.DAT:\*
- 17: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1996.DAT:\*
- 18: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1997.DAT:\*
- 19: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1998.DAT:\*
- 20: /SID2/gcgdata/geneseq/geneseqp-emb1/AA1999.DAT:\*
- 21: /SID2/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT:\*
- 22: /SID2/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT:\*
- 23: /SID2/gcgdata/geneseq/geneseqp-emb1/AA2002.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	181	91.0	419	22	AAE08627
2	181	91.0	419	22	AAE08628
3	181	91.0	419	22	AAE08629
4	181	91.0	419	22	AAE08627
5	181	91.0	419	22	AAE08627
6	173	86.9	44	20	AAV18297
7	173	86.9	44	20	AAV18300
8	173	86.9	44	20	AAV18301
9	173	86.9	419	22	AAE08630
10	173	86.9	419	22	AAE08630

11	173	86.9	419	22	AAE08628	Human protein C de
12	169	84.9	44	20	AAV18303	Human protein C de
13	169	84.9	44	22	AAE08627	Human protein C de
14	169	84.9	45	19	AAV75710	Human protein C de
15	169	84.9	415	21	AAV56803	Human protein C de
16	169	84.9	419	14	AAV72753	Human protein C de
17	169	84.9	419	19	AAV72753	Human protein C de
18	169	84.9	419	22	AAE08625	Human protein C de
19	169	84.9	419	22	AAE08627	Human protein C de
20	169	84.9	419	22	AAE08628	Human protein C de
21	169	84.9	419	22	AAE08629	Human protein C de
22	169	84.9	419	22	AAE08627	Human protein C de
23	169	84.9	419	22	AAE08628	Human protein C de
24	169	84.9	419	23	AAU99002	Human protein C de
25	169	84.9	419	23	AAU99003	Human protein C de
26	169	84.9	419	23	AAU99004	Human protein C de
27	169	84.9	419	23	AAU99005	Human protein C de
28	169	84.9	419	23	AAU99006	Human protein C de
29	169	84.9	419	23	AAU99007	Human protein C de
30	169	84.9	419	23	AAU99008	Human protein C de
31	169	84.9	419	23	AAU99009	Human protein C de
32	169	84.9	419	23	AAU99010	Human protein C de
33	169	84.9	419	23	AAU99011	Human protein C de
34	169	84.9	419	23	AAU99012	Human protein C de
35	169	84.9	419	23	AAU99013	Human protein C de
36	169	84.9	419	23	AAU99014	Human protein C de
37	169	84.9	419	23	AAU99015	Human protein C de
38	169	84.9	419	23	AAU99016	Human protein C de
39	169	84.9	419	23	AAU99017	Human protein C de
40	169	84.9	419	23	AAU99018	Human protein C de
41	169	84.9	419	23	AAU99019	Human protein C de
42	169	84.9	419	23	AAU99020	Human protein C de
43	169	84.9	419	23	AAU99021	Human protein C de
44	169	84.9	419	23	AAU99022	Human protein C de
45	169	84.9	419	23	AAU99023	Human protein C de

#### ALIGNMENTS

RESULT 1	AAE08627	standard; Protein: 419 AA.
ID	AAE08627	standard; Protein: 419 AA.
AC	AAE08627	
XX	01-NOV-2001	(first entry)
DT	01-NOV-2001	(first entry)
DE	Human protein C derivative #1.	
XX	Human, protein C derivative; antileukemia activity; thrombosis;	
XX	serpin inactivation; acute coronary syndrome; myocardial infarction;	
XX	vascular occlusive disorder; hypercoagulable state; angina; sepsis;	
XX	disseminated intravascular coagulation; DIC; burn; transplantation;	
XX	sickle cell disease; viral haemorrhagic fever; protein C deficiency;	
XX	haemolytic uraemic syndrome; acute arterial thrombotic occlusion;	
XX	thromboembolism; prothrombotic disorder; gene therapy; thalassemia.	
OS	Homo sapiens.	
XX	WO200159084-A1.	
PN	16-AUG-2001.	
PD	02-FEB-2001; 2001WO-US01221.	
PF	11-FEB-2000; 2000US-0181948.	
PR	14-MAR-2000; 2000US-0189199.	
XX	(ELIL ) LILLY & CO ELI.	
PA	Gerlitz BE, Grinnell BW, Jones BE;	
XX		
PI		
XX		

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DR WPI: 2001-514662/56.
XR N-PSDB: AAD15225.
PT Protein C derivative for treating acute coronary syndromes, vascular
PT occlusive disorders, thrombotic disorders and sepsis, comprises
PT substitutions at specified amino acid positions
PS Claim 3; Page 46-47; 59pp; English.
XX
XX The invention relates to human protein C derivatives and nucleic acid
XX molecules encoding such derivatives. These derivatives have increased
XX anticoagulation activity, resistance to serpin inactivation and
XX increased sensitivity to thrombin activation compared to wild type
XX protein C, and retains the biological activity of the wild type human
XX protein C. Protein C derivatives are useful in the manufacture of a
XX medicament for the treatment of acute coronary syndromes e.g. myocardial
XX infarction and unstable angina; and disease states predisposing to
XX thrombosis; vascular occlusive disorders and hypercoagulable states e.g.
XX disseminated intravascular coagulation (DIC), burns, transplantations,
XX thalassaemia, sickle cell disease, viral haemorrhagic fever and
XX haemolytic uremic syndrome; sepsis in combination with bacterial
XX permeability increasing protein; thrombotic disorders in combination
XX with an anti-platelet agent; protein C deficiency; acute arterial
XX thrombotic occlusion, thromboembolism or stenosis in coronary, cerebral
XX or peripheral arteries or in vascular grafts in combination with a
XX thrombolytic agent. Nucleic acid molecules of the invention are useful
XX for treating humans with genetically predisposed prothrombotic disorders
XX by gene therapy. The present sequence is human protein C derivative.
XX
SQ Sequence 419 AA;
Query Match 91.0%; Score 181; DB 22; Length 419;
Best Local Similarity 77.3%; Pred. No. 1.1e-21;
Matches 34; Conservative 1; Mismatches 9; Indels 0; Gaps 0
OY 1 ANSFLXLRHGSILRXCIIXICDFXXAKXIFEDVDDTLAFWSKH 44
||||| ||||||| || |||| || ||||| |||||||
Db 1 ANSFLLELRHGSILRERCIEICDFEAKKIFEDVDTLFWSKH 44
RESULT 2
AAE08628
ID AAE08628 standard; Protein: 419 AA.
XX
XX AAE08628;
XX
XX DT 01-NOV-2001 (first entry)
XX
XX DE Human protein C derivative #2.
XX
XX Human; protein C derivative; anticoagulation activity; thrombosis;
XX serpin inactivation; acute coronary syndrome; myocardial infarction;
XX vascular occlusive disorder; hypercoagulable state; angina; sepsis;
XX disseminated intravascular coagulation; DIC; burn; transplantation;
XX sickle cell disease; viral haemorrhagic fever; protein C deficiency;
XX haemolytic uremic syndrome; acute arterial thrombotic occlusion;
XX thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.
XX
XX OS Homo sapiens.
XX
XX PN WO200159084-A1.
XX
XX PD 16-AUG-2001.
XX
XX PF 02-FEB-2001; 2001WO-US01221.
XX
XX PR 11-FEB-2000; 2000US-0181948.
XX
XX PR 14-MAR-2000; 2000US-0189199.
XX
XX PA (ELIT ) LILLY & CO ELI.
XX
XX Gerlitz BE, Grinnell BW, Jones BE.
XX

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DR	WP1: 2001-514662/56.
DR	N-PSDB; AAD15226.
XX	Protein C derivative for treating acute coronary syndromes, vascular
PT	occlusive disorders, thrombotic disorders and sepsis, comprises
PT	substitutions at specified amino acid positions
PS	Claim 4; Page 47-48; 59pp; English.
XX	The invention relates to human protein C derivatives and nucleic acid
CC	molecules encoding such derivatives. These derivatives have increased
CC	anticoagulation activity, resistance to serpin inactivation and
CC	increased sensitivity to thrombin activation compared to wild type
CC	protein C, and retains the biological activity of the wild type human
CC	protein C. Protein C derivatives are useful in the manufacture of a
CC	medicament for the treatment of acute coronary syndromes e.g. myocardial
CC	infarction and unstable angina; and disease states predisposing to
CC	thrombosis; vascular occlusive disorders and hypercoagulable states e.g.
CC	dissminated intravascular coagulation (DIC), burns, transplantations,
CC	thalassaemia, sickle cell disease, viral haemorrhagic fever and
CC	haemolytic uremic syndrome; sepsis in combination with bacterial
CC	permeability increasing protein; thrombotic disorders in combination
CC	with an anti-platelet agent; protein C deficiency; acute arterial
CC	thrombotic occlusion, thromboembolism or stenosis in coronary, cerebral
CC	or peripheral arteries or in vascular grafts in combination with a
CC	thrombolytic agent. Nucleic acid molecules of the invention are useful
CC	for treating humans with genetically predisposed prothrombotic disorders
CC	by gene therapy. The present sequence is human protein C derivative.
XX	
SQ	Sequence 419 AA:
	Query Match 91.0%; Score 181; DB 22; Length 419;
	Best Local Similarity 77.3%; Pred. No. 1,1e-21;
	Matches 34; Conservative 1; Mismatches 9; Indels 0; Gaps 0.
OY	1 ANSFLXLRHGSILRXXCIXXICDPFXAKXIIFZVDVDTLAFMSKH 44                        :
Db	1 ANSFLEELRHGSLERECIEICDFEAKEIFEDVDTLAFMSKH 44
RESULT 3	
AAE08629	ID AAE08629 standard; Protein; 419 AA.
XX	AAE08629;
XX	
DT	01-NOV-2001 (first entry)
XX	
DE	Human protein C derivative #3.
XX	
KW	Human; protein C derivative; anticoagulation activity; thrombosis;
KW	serpin inactivation; acute coronary syndrome; myocardial infarction;
KW	vascular occlusive disorder; hypercoagulable state; angina; sepsis;
KW	dissminated intravascular coagulation; DIC; burn; transplantation;
KW	sickle cell disease; viral haemorrhagic fever; protein C deficiency;
KW	haemolytic uremic syndrome; acute arterial thrombotic occlusion;
KW	thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.
XX	
OS	Homo sapiens.
XX	
FH	Key Location/Qualifiers
FT	Misc-difference 10
FT	/note= "Encoded by CA"
XX	
PN	W0200159084-A1.
XX	
PD	16-AUG-2001.
XX	
PF	02-FEB-2001; 2001MO-US01221.
XX	
PR	11-FEB-2000; 2000US-0181948.
XX	
PR	14-MAR-2000; 2000US-0189199.
XX	

PA (ELIL ) LILLY & CO ELI.  
 XX Gerlitz BE, Grinnell BW, Jones BE.  
 XX WPI, 2001-514662/56.  
 DR N-PSDB; AAD15227.  
 XX  
 PT Protein C derivative for treating acute coronary syndromes, vascular  
 PT occlusive disorders, thrombotic disorders and sepsis, comprises  
 PT substitutions at specified amino acid positions -  
 XX  
 XX Claim 5; Page 48-49; 59pp: English.  
 XX  
 PS The invention relates to human protein C derivatives and nucleic acid  
 CC molecules encoding such derivatives. These derivatives have increased  
 CC anticoagulation activity, resistance to serpin inactivation and  
 CC increased sensitivity to thrombin activation compared to wild type  
 CC protein C, and retains the biological activity of the wild type human  
 CC protein C. Protein C derivatives are useful in the manufacture of a  
 CC medicament for the treatment of acute coronary syndromes e.g. myocardial  
 CC infarction and unstable angina; and disease states predisposing to  
 CC thrombosis; vascular occlusive disorders and hypercoagulable states e.g.  
 CC disseminated intravascular coagulation (DIC), burns, transplantations,  
 CC cholesteraemia, sickle cell disease, viral haemorrhagic fever and  
 CC haemolytic uremic syndrome; sepsis in combination with bacterial  
 CC permeability increasing protein; thrombotic disorders in combination  
 CC with an anti-platelet agent; protein C deficiency; acute arterial  
 CC thrombotic occlusion, thromboembolism or stenosis in coronary, cerebral  
 CC or peripheral arteries or in vascular grafts in combination with a  
 CC thrombolytic agent. Nucleic acid molecules of the invention are useful  
 CC for treating humans with genetically predisposed prothrombotic disorders  
 CC by gene therapy. The present sequence is human protein C derivative.  
 XX

SO Sequence 419 AA:

Query Match 91.0%; Score 181; DB 22; Length 419;  
 Best Local Similarity 77.3%; Pred. No. 1.1e-21;  
 Matches 34; Conservative 1; Mismatches 9; Indels 0; Gaps 0;

OY 1 ANSEFLXXLRHGSIXRCIXXICDPXAXKXFEZVDOTLAFMSKH 44  
 1 ANSFLELRHGSLERECTERICDFEAKELFEDVDTLAFMSKH 44

RESULT 4  
 AAB82675  
 ID AAB82675 standard; Protein; 419 AA.

AC AAB82675;

DT 15-OCT-2001 (first entry)

DE Human protein C derivative (S11G/Q32E/N33D/L194S).

XX  
 XX protein C; human; coronary syndrome; thrombosis; angina;  
 KM myocardial infarction; vascular occlusive disorder;  
 KM hypercoagulation; sepsis; protein C deficiency; occlusion;  
 KM thromboembolism; stenosis; antibacterial; immunosuppressive;  
 KM thrombolytic; cardiac; antiangiinal; anticoagulant; therapy;  
 mutant; mutain.  
 XX  
 XX

OS Homo saplens.

OS Synthetic.

XX  
 FH Key Location/Qualifiers

FT MISC-difference 11 /note= "Ser in wild-type protein"

FT MISC-difference 32 /note= "Gln in wild-type protein"

FT MISC-difference 33 /note= "Asn in wild-type protein"

FT MISC-difference 194 /note= "Leu in wild-type protein"

FT Domain 1..45  
 FT /note= "Gla. domain"  
 FT Disulfide-bond 50..69  
 FT Disulfide-bond 59..64  
 FT Disulfide-bond 80..89  
 FT Disulfide-bond 98..109  
 FT Disulfide-bond 120..133  
 FT Disulfide-bond 141..277  
 FT Disulfide-bond 146..212  
 FT Disulfide-bond 331..345  
 FT Disulfide-bond 356..384  
 FT Disulfide-bond 156..157  
 FT /note= "Cleaveage makes a 2-chain inactive  
 FT precursor (155-amino acid light chain  
 FT attached via a disulfide bond to a  
 FT 262-amino acid heavy chain)"  
 FT  
 FT Modified-site 6  
 FT /note= "gamma-carboxylated"  
 FT Modified-site 7  
 FT /note= "gamma-carboxylated"  
 FT Modified-site 14  
 FT /note= "gamma-carboxylated"  
 FT Modified-site 16  
 FT /note= "gamma-carboxylated"  
 FT Modified-site 19  
 FT /note= "gamma-carboxylated"  
 FT Modified-site 20  
 FT /note= "gamma-carboxylated"  
 FT Modified-site 25  
 FT /note= "gamma-carboxylated"  
 FT Modified-site 26  
 FT /note= "gamma-carboxylated"  
 FT Peptide 158..169  
 FT /note= "activation peptide; removal activates the  
 FT 2-chain zymogen"  
 FT  
 FT Cleavage-site 169..170  
 FT /note= "chrombin cleavage site"  
 FT Modified-site 29  
 FT /note= "N-glycosylated"  
 FT Modified-site 248  
 FT /note= "N-glycosylated"  
 FT Modified-site 313  
 FT /note= "N-glycosylated"  
 FT Modified-site 329  
 FT /note= "N-glycosylated"  
 FT  
 XX WO200157193-A2.  
 PN  
 XX  
 PD 09-AUG-2001.  
 XX  
 PD 19-JAN-2001; 2001WO-US00020.  
 XX  
 XX  
 PR 02-FEB-2000; 2000US-0179801.  
 PR 14-MAR-2000; 2000US-0189197.  
 XX  
 PA (ELIL ) LILLY & CO ELI.  
 XX  
 XX Gerlitz BE, Jones BE.  
 PT WPI, 2001-496919/54.  
 DR N-PSDB; AAB26363.  
 XX  
 PT Novel human protein C derivative for treating, e.g., myocardial  
 PT infarction, unstable angina, sepsis, thrombotic disorders, acute  
 PT arterial thrombotic occlusion, and thromboembolism -  
 XX  
 PS Claim 3; Page 52-53; 63pp: English.  
 XX  
 CC The present sequence is that of a claimed human protein C  
 CC derivative in which Ser at amino acid position 11 of the mature  
 CC wild-type protein C sequence (see AAB82673) is substituted with  
 CC Gly, Gln at position 32 with Glu, Asn at position 33 with Asp, and  
 CC Leu at position 194 with Ser. The protein is an example of protein

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CC C derivatives of the invention that have at least 2 amino acid
CC substitutions, but which have increased anticoagulant activity and
CC resistance to inactivation by serpins compared with the wild-type
CC protein, while retaining the biological activity of the wild-type
CC protein. A method of producing the derivatives using recombinant
CC DNA methods is claimed. The protein C derivatives are useful for
CC treating coronary syndromes and disease states predisposing to
CC thrombosis (e.g. myocardial infarction and unstable angina),
CC vascular occlusive disorders and hypercoagulable states, sepsis (in
CC combination with bactericidal permeability increasing protein or
CC with tissue factor pathway inhibitor), thrombotic disorders (in
CC combination with an anti-platelet agent or by local delivery through
CC an intracoronary catheter), protein C deficiency, acute arterial
CC thrombotic occlusion, thromboembolism, or stenosis in coronary,
CC cerebral or peripheral arteries or in vascular grafts. Human
CC patients with genetically predisposed prothrombotic disorders may
CC be treated by gene therapy (all claimed).
XX
SQ Sequence 419 AA:

Query Match 91.0%; Score 181; DB 22; Length 419;
Best Local Similarity 77.3%; Pred. No. 1.1e-21;
Matches 34; Conservative 1; Mismatches 9; Indels 0; Gaps 0:

Oy 1 ANSFLXXLRHSGSLRXKXIXICDFXXAKXIFZDDVDTLAFWSKH 44
    ||||| ||||| || ||||| ||:|||||||
Db 1 ANSFLLELRHSGSLRRCIEICDFEAKEIFEDVDTLAFWSKH 44

RESULT 5
AAB82676
ID AAB82676 standard; Protein: 419 AA.
XX
AC AAB82676;
XX
DT 15-OCT-2001 (first entry)
XX
DE Human protein C derivative (S11G/Q32E/N33D/L194S/T754S).
XX
KW Protein C; human; coronary syndrome; thrombosis; angina;
KW myocardial infarction; vascular occlusive disorder;
KW hypercoagulation; sepsis; protein C deficiency; occlusion;
KW thromboembolism; stenosis; antibacterial; immunosuppressive;
KW thrombolytic; cardiact; antiangiinal; anticoagulant; therapy;
KW mutant; mutein.
XX
OS Homo sapiens.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Misc-difference 11 /note= "Ser in wild-type protein"
FT Misc-difference 32 /note= "Gln in wild-type protein"
FT Misc-difference 33 /note= "Asn in wild-type protein"
FT Misc-difference 194 /note= "Leu in wild-type protein"
FT Misc-difference 254 /note= "Thr in wild-type protein"
FT Domain 1..45 /note= "G1a domain"
FT Disulfide-bond 50..69 /note= "G1a domain"
FT Disulfide-bond 59..64
FT Disulfide-bond 80..89
FT Disulfide-bond 98..109
FT Disulfide-bond 120..133
FT Disulfide-bond 141..277
FT Disulfide-bond 196..212
FT Disulfide-bond 331..345
FT Disulfide-bond 356..384
FT Disulfide-bond 156..157
FT /note= "cleavage makes a 2-chain inactive

```

```

FT FT precursor (155-amino acid light chain
FT FT attached via a disulfide bond to a
FT FT 262-amino acid heavy chain)"
FT FT
FT FT Modified-site 6 /note= "gamma-carboxylated"
FT FT Modified-site 7 /note= "gamma-carboxylated"
FT FT Modified-site 14 /note= "gamma-carboxylated"
FT FT Modified-site 16 /note= "gamma-carboxylated"
FT FT Modified-site 19 /note= "gamma-carboxylated"
FT FT Modified-site 20 /note= "gamma-carboxylated"
FT FT Modified-site 25 /note= "gamma-carboxylated"
FT FT Modified-site 26 /note= "gamma-carboxylated"
FT FT Peptide 158..169 /note= "activation peptide; removal activates the
FT FT 2-chain zymogen"
FT FT Cleavage-site 169..170 /note= "thrombin cleavage site"
FT FT Modified-site 29 /note= "N-glycosylated"
FT FT Modified-site 248 /note= "N-glycosylated"
FT FT Modified-site 313 /note= "N-glycosylated"
FT FT Modified-site 329 /note= "N-glycosylated"
FT FT Modified-site /note= "N-glycosylated"
FT FT
FT FT WO200157193-A2.
FT FT
FT FT 09-AUG-2001.
FT FT
FT FT 19-JAN-2001; 2001WO-US00020.
FT FT
FT FT 02-FEB-2000; 2000US-0179801.
FT FT 14-MAR-2000; 2000US-0189197.
FT FT
FT FT (ELLI ) LILLY & CO ELLI.
FT FT
FT FT Gerlitz BE, Jones BE;
FT FT
FT FT WPI; 2001-496919/54.
FT FT N-PSDB; AAH26364.
FT FT
FT FT Novel human protein C derivative for treating, e.g., myocardial
FT FT infarction, unstable angina, sepsis, thrombotic disorders, acute
FT FT arterial thrombotic occlusion, and thromboembolism -
XX
XX
XX Claim 4; Page 53-54; 63pp; English.
XX
XX The present sequence is that of a claimed human protein C derivative
XX in which Ser at position 11 of the mature wild-type protein C
XX sequence (see AAB82673) is substituted with Gly, Gln at position 32
XX with Gln, Asn at position 33 with Asp, Leu at position 194 with Ser,
XX and Thr at position 254 with Ser. It is an example of protein C
XX derivatives of the invention that have at least 2 amino acid
XX substitutions, but which have increased anticoagulant activity and
XX resistance to inactivation by serpins compared with the wild-type
XX protein, while retaining the biological activity of the wild-type
XX protein. A method of producing the derivatives using recombinant
XX DNA methods is claimed. The protein C derivatives are useful for
XX treating coronary syndromes and disease states predisposing to
XX thrombosis (e.g. myocardial infarction and unstable angina),
XX vascular occlusive disorders and hypercoagulable states, sepsis (in
XX combination with bactericidal permeability increasing protein or
XX with tissue factor pathway inhibitor), thrombotic disorders (in
XX combination with an anti-platelet agent or by local delivery through
XX an intracoronary catheter), protein C deficiency, acute arterial

```

CC thrombotic occlusion, thromboembolism, or stenosis in coronary,  
 CC cerebral or peripheral arteries or in vascular grafts. Human  
 CC patients with genetically predisposed prothrombotic disorders may  
 CC be treated by gene therapy (all claimed).  
 XX

SO Sequence 419 AA:

Query Match 91.0%; Score 181; DB 22; Length 419;  
 Best Local Similarity 77.3%; Pred. No. 1.1e-21;  
 Matches 34; Conservative 1; Mismatches 9; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFZDVDDTLAFWSKH 44  
 DB 1 ANSFLXELRHGSLRRCIEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 6

AAI18297  
 ID AAY18297 standard; peptide: 44 AA.

AC AAY18297;

DT 17-AUG-1999 (first entry)

DE Modified GLA domain of vitamin K-dependent protein.

KW GLA domain; muteln; vitamin K-dependent protein; clotting disorder;  
 therapy.

OS Homo sapiens.  
 OS Synthetic.

Key Location/Qualifiers

FT Misc-difference 1..44 /note="Xaa=gamma-carboxyglutamic acid, or glutamic  
 FT acid"

XX WO9920767-A1.

XX 29-APR-1999.

XX 20-OCT-1998; 98WO-US22152.

XX 23-OCT-1997; 97US-0955636.

XX (MINU ) UNIV MINNESOTA.

XX Nelstuen GL;

XX WPI: 1999-288309/24.

XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic  
 XX acid domain, useful for treating clotting disorders

XX Claim 6: Page 78; 86pp; English.

XX This sequence represents a modified GLA (gamma-carboxyglutamic acid)  
 CC domain. The invention relates to a vitamin K-dependent polypeptide  
 CC comprising a modified GLA domain containing an amino acid substitution  
 CC which enhances membrane binding of the modified polypeptide as compared  
 CC to the native polypeptide. The polypeptide is used to treat a clotting  
 CC disorder by decreasing or increasing clot formation. Modification of the  
 CC GLA domain results in a protein which has enhanced membrane binding  
 CC affinity as compared to the native protein.  
 XX

SO Sequence 44 AA:

Query Match 87.9%; Score 175; DB 20; Length 44;

Best Local Similarity 95.5%; Pred. No. 1e-21;  
 Matches 42; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFZDVDDTLAFWSKH 44  
 XXX

DB 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFEDVDDTLAFWSKH 44

RESULT 7

AAI18300  
 ID AAY18300 standard; peptide: 44 AA.

AC AAY18300;

DT 17-AUG-1999 (first entry)

DE Modified GLA domain of vitamin K-dependent protein.

KW GLA domain; muteln; vitamin K-dependent protein; clotting disorder;  
 therapy.

OS Homo sapiens.  
 OS Synthetic.

Key Location/Qualifiers

FT Misc-difference 1..44 /note="Xaa=gamma-carboxyglutamic acid, or glutamic  
 FT acid"

XX WO9920767-A1.

XX 29-APR-1999.

XX 20-OCT-1998; 98WO-US22152.

XX 23-OCT-1997; 97US-0955636.

XX (MINU ) UNIV MINNESOTA.

XX Nelstuen GL;

XX WPI: 1999-288309/24.

XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic  
 XX acid domain, useful for treating clotting disorders

XX Claim 9: Page 79; 86pp; English.

XX This sequence represents a modified GLA (gamma-carboxyglutamic acid)  
 CC domain. The invention relates to a vitamin K-dependent polypeptide  
 CC comprising a modified GLA domain containing an amino acid substitution  
 CC which enhances membrane binding of the modified polypeptide as compared  
 CC to the native polypeptide. The polypeptide is used to treat a clotting  
 CC disorder by decreasing or increasing clot formation. Modification of the  
 CC GLA domain results in a protein which has enhanced membrane binding  
 CC affinity as compared to the native protein.  
 XX

SO Sequence 44 AA:

Query Match 86.9%; Score 173; DB 20; Length 44;  
 Best Local Similarity 95.5%; Pred. No. 2.1e-21;  
 Matches 42; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFZDVDDTLAFWSKH 44  
 DB 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFEDVDDTLAFWSKH 44

RESULT 8

AAI18301  
 ID AAY18301 standard; peptide: 44 AA.

AC AAY18301;

DT 17-AUG-1999 (first entry)

XX Modified GLA domain of vitamin K-dependent protein.  
 XX

```

KM GLA domain; mutein; vitamin K-dependent protein; clotting disorder;
KW therapy.
XX
XX Homo sapiens.
OS Synthetic.
OS
XX Key Location/Qualifiers
FH Misc-difference 1..44
FT /note= "Xaa= gamma-carboxyglutamic acid, or glutamic
FT acid"
XX
XX WO9920767-A1.
XX
XX 29-APR-1999.
XX
XX 20-OCT-1998; 98WO-US22152.
XX
XX 23-OCT-1997; 97US-0955636.
XX
XX (MINU ) UNIV MINNESOTA.
XX
XX Nelstuen GL.
XX
XX WPI, 1999-288309/24.
XX
XX Vitamin K-dependent polypeptide with modified gamma-carboxyglutamic
XX acid domain, useful for treating clotting disorders
XX
XX Claim 9; Page 82; 86pp; English.
XX
XX This sequence represents a modified GLA (gamma-carboxyglutamic acid)
XX domain. The invention relates to a vitamin K-dependent polypeptide
XX comprising a modified GLA domain containing an amino acid substitution
XX which enhances membrane binding of the modified polypeptide as compared
XX to the native polypeptide. The polypeptide is used to treat a clotting
XX disorder by decreasing or increasing clot formation. Modification of the
XX GLA domain results in a protein which has enhanced membrane binding
XX affinity as compared to the native protein.
XX
XX SQ Sequence 44 AA;
XX
XX Query Match 86.9%; Score 173; DB 20; Length 44;
XX Best Local Similarity 95.5%; Pred. No. 2.1e-21;
XX Matches 42; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
XX
XX OY 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFZDVDDTLAFWSKH 44
XX ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFZDVDDTLAFWSKH 44
XX
XX Db
XX
XX RESULT 9
XX AAE08630
XX ID AAE08630 standard; Protein: 419 AA.
XX
XX AC AAE08630;
XX
XX 01-NOV-2001 (first entry)
XX
XX Human protein C derivative #4.
XX
XX Human; protein C derivative; anticoagulation activity; thrombosis;
XX serpin inactivation; acute coronary syndrome; myocardial infarction;
XX vascular occlusive disorder; hypercoagulable state; angina; sepsis;
XX disseminated intravascular coagulation; DIC; burn; transplantation;
XX sickle cell disease; viral haemorrhagic fever; protein C deficiency;
XX haemolytic uremic syndrome; acute arterial thrombotic occlusion;
XX thromboembolism; prothrombotic disorder; gene therapy; thalassaemia.
XX
XX Homo sapiens.
XX
XX OS Synthetic.
XX
XX Key Location/Qualifiers
XX Misc-difference 10
XX /note= "His in wild-type protein"
XX
XX PD 16-AUG-2001.

```

```

XX
XX 02-FEB-2001; 2001WO-US01221.
XX
XX 11-FEB-2000; 2000US-0181948.
XX
XX 14-MAR-2000; 2000US-0189199.
XX
XX (ELIT ) LILLY & CO ELI.
XX
XX Gerlitz BE, Grinnell BW, Jones BE;
XX
XX WPI, 2001-514662/56.
XX
XX N-PSDB; AAD15228.
XX
XX Protein C derivative for treating acute coronary syndromes, vascular
XX occlusive disorders, thrombotic disorders and sepsis, comprises
XX substitutions at specified amino acid positions .
XX
XX Claim 6; Page 50-51; 59pp; English.
XX
XX The invention relates to human protein C derivatives and nucleic acid
XX molecules encoding such derivatives. These derivatives have increased
XX anticoagulation activity, resistance to serpin inactivation and
XX increased sensitivity to thrombin activation compared to wild type
XX protein C, and retains the biological activity of the wild type human
XX protein C. Protein C derivatives are useful in the manufacture of a
XX medicament for the treatment of acute coronary syndromes e.g. myocardial
XX infarction and unstable angina; and disease states predisposing to
XX thrombosis; vascular occlusive disorders and hypercoagulable states e.g.
XX disseminated intravascular coagulation (DIC), burns, transplantations,
XX thalassaemia, sickle cell disease, viral haemorrhagic fever and
XX haemolytic uremic syndrome; sepsis in combination with bacterial
XX permeability increasing protein; thrombotic disorders in combination
XX with an anti-platelet agent; protein C deficiency; acute arterial
XX thrombotic occlusion, thromboembolism or stenosis in coronary, cerebral
XX or peripheral arteries or in vascular grafts in combination with a
XX thrombolytic agent. Nucleic acid molecules of the invention are useful
XX for treating humans with genetically predisposed prothrombotic disorders
XX by gene therapy. The present sequence is human protein C derivative.
XX
XX SQ Sequence 419 AA;
XX
XX Query Match 86.9%; Score 173; DB 22; Length 419;
XX Best Local Similarity 75.0%; Pred. No. 2.3e-20;
XX Matches 33; Conservative 1; Mismatches 10; Indels 0; Gaps 0;
XX
XX OY 1 ANSFLXXLRHGSIXRCIXXICDPFXAKXIFZDVDDTLAFWSKH 44
XX ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
XX 1 ANSFLXELRGSLERCIERICDPFEAKELFEDVDTLAFWSKH 44
XX
XX Db
XX
XX RESULT 10
XX AAB82677
XX ID AAB82677 standard; Protein: 419 AA.
XX
XX AC AAB82677;
XX
XX 15-OCT-2001 (first entry)
XX
XX Human protein C derivative (H10Q/S11G/O32E/N33D/LJ194S).
XX
XX Protein C; human; coronary syndrome; thrombosis; angina;
XX myocardial infarction; vascular occlusive disorder;
XX hypercoagulable state; sepsis; protein C deficiency; occlusion;
XX thromboembolism; stenosis; antibacterial; immunosuppressive;
XX thrombolytic; cardiant; antianginal; anticoagulant; therapy;
XX mutant; mutein.
XX
XX Homo sapiens.
XX
XX OS Synthetic.
XX
XX Key Location/Qualifiers
XX Misc-difference 10
XX /note= "His in wild-type protein"
XX
XX FT

```

```

FT Misc-difference 11 /note= "Ser in wild-type protein"
FT Misc-difference 32 /note= "Gln in wild-type protein"
FT Misc-difference 33 /note= "Asn in wild-type protein"
FT Misc-difference 194 /note= "Leu in wild-type protein"
FT Domain 1..45 /note= "Gla domain"
FT Disulfide-bond 50..69
FT Disulfide-bond 59..64
FT Disulfide-bond 80..89
FT Disulfide-bond 98..109
FT Disulfide-bond 120..133
FT Disulfide-bond 141..277
FT Disulfide-bond 196..212
FT Disulfide-bond 331..345
FT Disulfide-bond 356..384
FT Cleavage-site 156..157 /note= "cleavage makes a 2-chain inactive precursor (155-amino acid light chain attached via a disulfide bond to a 262-amino acid heavy chain)"
FT Modified-site 6 /note= "gamma-carboxylated"
FT Modified-site 7 /note= "gamma-carboxylated"
FT Modified-site 14 /note= "gamma-carboxylated"
FT Modified-site 16 /note= "gamma-carboxylated"
FT Modified-site 19 /note= "gamma-carboxylated"
FT Modified-site 20 /note= "gamma-carboxylated"
FT Modified-site 25 /note= "gamma-carboxylated"
FT Modified-site 26 /note= "gamma-carboxylated"
FT Modified-site /note= "gamma-carboxylated"
FT Peptide 158..169 /note= "activation peptide; removal activates the 2-chain zymogen"
FT Cleavage-site 169..170
FT Modified-site /note= "thrombin cleavage site"
FT Modified-site 29 /note= "N-glycosylated"
FT Modified-site 248 /note= "N-glycosylated"
FT Modified-site 313 /note= "N-glycosylated"
FT Modified-site 329 /note= "N-glycosylated"
FT Modified-site /note= "N-glycosylated"
FT WO200157193-A2.
FT 09-AUG-2001.
FT PD 19-JAN-2001: 2001WO-US00020.
FT XX 02-FEB-2000: 2000US-0179801.
FT PR 14-MAR-2000: 2000US-0189197.
FT XX (ELIT ) LILLY & CO ELI.
FT PA Gerlitz BE, Jones BE;
FT PI WPI: 2001-496919/54.
FT DR N-PSDB: AAI26365.
FT XX Novel human protein C derivative for treating, e.g., myocardial
FT PT infarction, unstable angina, sepsis, thrombotic disorders, acute
FT PT arterial thrombotic occlusion, and thromboembolism -

```

```

XX Claim 5; Page 54-55; 63pp; English.
PS The present sequence is that of a claimed human protein C derivative
XX in which His at position 10 of the mature wild-type protein C
CC sequence (see AAB82673) is substituted with Gln, Ser at position 11
CC with Gly, Gln at position 32 with Gln, Asn at position 33 with Asp,
CC and Leu at position 194 with Ser. It is an example of protein C
CC derivatives of the invention that have at least 2 amino acid
CC substitutions, but which have increased anticoagulant activity and
CC resistance to inactivation by serpins compared with the wild-type
CC protein. A method of producing the derivatives using recombinant
CC DNA methods is claimed. The protein C derivatives are useful for
CC treating coronary syndromes and disease states predisposing to
CC thrombosis (e.g. myocardial infarction and unstable angina),
CC vascular occlusive disorders and hypercoagulable states, sepsis (in
CC combination with bactericidal permeability increasing protein or
CC with tissue factor pathway inhibitor), thrombotic disorders (in
CC combination with an anti-platelet agent or by local delivery through
CC an intracoronary catheter), protein C deficiency, acute arterial
CC thrombotic occlusion, thromboembolism, or stenosis in coronary,
CC cerebral or peripheral arteries or in vascular grafts. Human
CC patients with genetically predisposed prothrombotic disorders may
CC be treated by gene therapy (all claimed).
XX
SQ Sequence 419 AA:
Query Match 86.9%; Score 173; DB 22; Length 419;
Best Local Similarity 75.0%; Pred. No. 2,3e-20;
Matches 33; Conservative 1; Mismatches 10; Indels 0; Gaps 0;
OY 1 ANSEFLXXLRHGSIXRXICIXXICDFXXAKKIFZDVDDTLAFWSKH 44
DB 1 ANSFLELRGSLERBCIEBICDFEAKKEIFEDVDTLAFWSKH 44
RESULT 11
AAB82678
ID AAB82678 standard; Protein: 419 AA.
XX
XX AAB82678;
AC 15-OCT-2001 (first entry)
XX
XX Human protein C derivative (H10Q/S11G/Q32E/N33D/L194S/T254S).
DE Protein C; human; coronary syndrome; thrombosis; angina;
XX myocardial infarction; vascular occlusive disorder;
KW hypercoagulation; sepsis; protein C deficiency; occlusion;
KW thromboembolism; stenosis; antibacterial; immunosuppressive;
KW thrombolytic; cardiac; antiangiinal; anticoagulant; therapy;
KW mutant; mutlein.
XX
XX Homo sapiens.
OS Synthetic.
OS
XX
XX key Location/Qualifiers
FT Misc-difference 10 /note= "His in wild-type protein"
FT Misc-difference 11 /note= "Ser in wild-type protein"
FT Misc-difference 32 /note= "Gln in wild-type protein"
FT Misc-difference 33 /note= "Gln in wild-type protein"
FT Misc-difference 33 /note= "Asn in wild-type protein"
FT Misc-difference 194 /note= "Leu in wild-type protein"
FT Misc-difference 254 /note= "Thr in wild-type protein"
FT Domain 1..45 /note= "Gla domain"
FT Disulfide-bond 50..69

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FT	Disulfide-bond	59..64	
FT	Disulfide-bond	80..89	
FT	Disulfide-bond	98..109	
FT	Disulfide-bond	120..133	
FT	Disulfide-bond	141..277	
FT	Disulfide-bond	196..212	
FT	Disulfide-bond	331..345	
FT	Disulfide-bond	356..384	
FT	Cleavage-site	156..157	
FT	/note=	"cleavage makes a 2-chain inactive precursor (155-amino acid light chain attached via a disulfide bond to a 262-amino acid heavy chain)"	
FT	Modified-site	6	
FT	/note=	"gamma-carboxylated"	
FT	Modified-site	7	
FT	/note=	"gamma-carboxylated"	
FT	Modified-site	14	
FT	/note=	"gamma-carboxylated"	
FT	Modified-site	16	
FT	/note=	"gamma-carboxylated"	
FT	Modified-site	19	
FT	/note=	"gamma-carboxylated"	
FT	Modified-site	20	
FT	/note=	"gamma-carboxylated"	
FT	Modified-site	25	
FT	/note=	"gamma-carboxylated"	
FT	Modified-site	26	
FT	/note=	"gamma-carboxylated"	
FT	Peptide	158..169	
FT	/note=	"activation peptide; removal activates the 2-chain zymogen"	
FT	Cleavage-site	169..170	
FT	/note=	"thrombin cleavage site"	
FT	Modified-site	29	
FT	/note=	"N-glycosylated"	
FT	Modified-site	248	
FT	/note=	"N-glycosylated"	
FT	Modified-site	313	
FT	/note=	"N-glycosylated"	
FT	Modified-site	329	
FT	/note=	"N-glycosylated"	
PN	WO200157193-A2.		
XX	09-AUG-2001.		
XX	19-JAN-2001.	2001WO-US00020.	
PE	02-FEB-2000.	2000US-0179801.	
PR	14-MAR-2000.	2000US-0189197.	
XX	(ELIL ) LILLY & CO ELI.		
PA			
XX			
XX	Gerlitz BE, Jones BE.		
DR	WPI: 2001-496919/54.		
XX			
PT	Novel human protein C derivative for treating, e.g., myocardial infarction, unstable angina, sepsis, thrombotic disorders, acute arterial thrombotic occlusion, and thromboembolism -		
PT			
XX			
PS	Claim 6; Page 56-57; 63pp: English.		
XX			
CC	The present sequence is that of a claimed human protein C derivative		
CC	at position 10 of the wild-type protein C sequence (see		
CC	ABB82673) is substituted with Gln, Ser at position 11 with Gly, Gln		
CC	at position 32 with Glu, Asn at position 33 with Asp, Leu at position		
CC	194 with Ser, and Thr at position 254 with Ser. It is an example of		
CC	protein C derivatives of the invention that have at least 2 amino acid		
CC	substitutions, but which have increased anticoagulant activity and		
CC	resistance to inactivation by serpins compared with the wild-type		
CC	protein, while retaining the biological activity of the wild-type		

CC	protein. A method of producing the derivatives using recombinant DNA methods is claimed. The protein C derivatives are useful for treating coronary syndromes and disease states predisposing to thrombosis (e.g. myocardial infarction and unstable angina).
CC	vascular occlusive disorders and hypercoagulable states, sepsis (in combination with bactericidal permeability increasing protein or with tissue factor pathway inhibitor), thrombotic disorders (in combination with an anti-platelet agent or by local delivery through an intracoronary catheter), protein C deficiency, acute arterial thrombotic occlusion, thromboembolism, or stenosis in coronary, cerebral or peripheral arteries or in vascular grafts. Human patients with genetically predisposed prothrombotic disorders may be treated by gene therapy (all claimed).
CC	
XX	
SQ	Sequence    419 AA;
Query Match	86.9%; Score 173; DB 22; Length 419;
Best Local Similarity	75.0%; Pred. No. 2,3e-20;
Matches    33;	Conservative    1; Mismatches    10; Indels        0; Gaps         0;
Oy	1 ANSFLXLRHGSIXRXCIXICDFYXAAXIFZDVDDTLAFWSKH 44 . 
Db	1 ANSFLEELRGSGLERECIEICDFEAKEIFEDVDVDTLAFWSKH 44
RESULT 12	
AAV18303	
ID	AAV18303 standard; peptide: 44 AA.
AC	AAV18303;
DT	17-AUG-1999 (first entry)
XX	
DE	Human protein C GLA domain.
XX	
KW	GLA domain: vitamin K-dependent protein; clotting disorder; therapy.
OS	Homo sapiens.
FH	Key : Location/Qualifiers
Misc-difference 1..44	/note= "Xaa= gamma-carboxylutamic acid, or glutamic acid"
WT	
WO9920767-A1.	
PN	
PD	29-APR-1999.
XX	
PF	20-OCT-1998; 98WO-US22152.
PR	23-OCT-1997; 97US-0955636.
PA	(MINU ) UNIV MINNESOTA.
NeIsestuen GL:	
WI: 1999-288309/24.	
Vitamin K-dependent polypeptide with modified gamma-carboxylutamic acid domain, useful for treating clotting disorders	
Disclosure: Page 14; 86pp; English.	
This sequence is the protein C GLA (gamma-carboxylutamic acid) domain. The invention relates to a vitamin K-dependent polypeptide comprising a modified GLA domain containing an amino acid substitution which enhances membrane binding of the modified polypeptide as compared to the native polypeptide. The polypeptide is used to treat a clotting disorder by decreasing or increasing clot formation. Modification of the GLA domain results in a protein which has enhanced membrane binding affinity as compared to the native protein.	



```

SQ Sequence 44 AA:
Query Match 84.9%; Score 169; DB 20; Length 44;
Best Local Similarity 93.2%; Pred. No. 1e-20;
Matches 41; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44
1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44

DB 1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44
1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44

RESULT 13
AAB36402 standard; peptide: 44 AA.
XX
AC AAB36402;
XX
DT 27-FEB-2001 (first entry)
XX
DE Human protein C gamma-carboxyglutamic acid domain SEQ ID NO:1.
XX
KM Vitamin K-dependent protein: factor VII; protein C; Gla domain;
KM gamma-carboxyglutamic acid domain; factor IX; protein S; protein Z;
KM factor X; prothrombin; enhanced membrane binding affinity;
KM clot formation; thrombolytic; haemostatic; bleeding disorder;
KM thrombosis; clotting disorder; haemophilia A; haemophilia B;
KM liver disease.
XX
OS Homo sapiens.
XX
PN WO200066753-A2.
XX
PD 09-NOV-2000.
XX
PE 28-APR-2000; 2000WO-US11416.
XX
PR 29-APR-1999; 9905-0302239.
XX
PA (MINU) UNIV MINNESOTA.
XX
PI Nelsstuen GL:
XX
DR WPI: 2001-007226/O1.
XX
PT Novel vitamin K-dependent polypeptide useful for treating clotting
PT disorders such as thrombosis and hemophilia, comprises modified
PT gamma-carboxy glutamic acid domain that enhances membrane binding
PT affinity -
XX
PS Example 5; Page 42; 81pp; English.
XX
CC The present invention describes a vitamin K-dependent polypeptide (I)
CC comprising a modified gamma-carboxy glutamic acid (Gla) domain having
CC at least one amino acid substitution, that enhances membrane binding
CC affinity and the activity of the polypeptide relative to a corresponding
CC native vitamin K-dependent polypeptide and inhibits clot formation.
CC (I) can have thrombolytic and haemostatic activities, and can be used
CC as an inhibitor of clot formation. (I) is useful for decreasing clot
CC formation in a mammal, a factor VII or factor IX containing a modified
CC Gla domain is useful for increasing clot formation and for treating a
CC bleeding disorder, including thrombosis and clotting disorders such as
CC haemophilia A, haemophilia B and liver disease. The present sequence
CC represents a human protein C Gla domain sequence, given in the
CC exemplification of the present invention.
XX
SQ Sequence 44 AA:
Query Match 84.9%; Score 169; DB 22; Length 44;
Best Local Similarity 93.2%; Pred. No. 1e-20;
Matches 41; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44
1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44

```

```

DB 1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44
1 ANSFLXXLRHGSLSRXICIXXICDFXAKXIFZDVDTLAFWSKH 44

RESULT 14
AAW75710 standard; protein: 45 AA.
XX
AC AAW75710;
XX
DT 08-DEC-1998 (first entry)
XX
DE Partial human protein C amino acid sequence.
XX
KM Gamma carboxyglutamic acid; human protein C; Gla domain; chimera;
KM pRC/RSV; RSV-PC; amplification, PCR, primer; transfection; anticoagulant;
KM human 293 cell; Protein S; myocardial infarction; venous thrombosis;
KM disseminated intravascular coagulation; thromboembolic disease; lupus;
KM adult respiratory distress syndrome; factor V Leiden; stroke.
XX
OS Homo sapiens.
XX
FH Key location/Qualifiers
FT Misc-difference 6 /note="Gamma carboxyglutamic acid"
FT Misc-difference 7 /note="Gamma carboxyglutamic acid"
FT Misc-difference 7 /note="Gamma carboxyglutamic acid"
FT Misc-difference 14 /note="Gamma carboxyglutamic acid"
FT Misc-difference 16 /note="Gamma carboxyglutamic acid"
FT Misc-difference 19 /note="Gamma carboxyglutamic acid"
FT Misc-difference 19 /note="Gamma carboxyglutamic acid"
FT Misc-difference 20 /note="Gamma carboxyglutamic acid"
FT Misc-difference 25 /note="Gamma carboxyglutamic acid"
FT Misc-difference 26 /note="Gamma carboxyglutamic acid"
FT Misc-difference 26 /note="Gamma carboxyglutamic acid"
FT Misc-difference 29 /note="Gamma carboxyglutamic acid"
XX
PN WO9820118-A1.
XX
PD 14-MAY-1998.
XX
PE 07-NOV-1997; 97WO-US20376.
XX
PR 25-JUL-1997; 97US-0053768.
XX
PR 08-NOV-1996; 96US-0745254.
XX
PA (OKLA-) OKLAHOMA MEDICAL RES FOUND.
XX
PI Esmon CT, Smirnov M;
XX
DR WPI: 1998-286934/25.
XX
PT Protein C chimeric proteins for use as anticoagulants - having gamma
PT carboxyglutamic acid region replaced with vitamin K dependent
PT clotting factor e.g. prothrombin
XX
PS Example 1; Page 15; 42pp; English.
XX
CC The present sequence represents the first three exons of the human
CC protein C protein, which contains gamma carboxyglutamic acid modified
CC residues. This sequence was replaced with the corresponding regions of
CC modified human prothrombin (AAW75709), to create a protein C prothrombin
CC Gla domain chimera. To produce this chimera, the wild-type protein C
CC cDNA was ligated into pRC/RSV to form RSV-PC, and then was digested with
CC restriction enzymes to remove the first three exons and the first codon
CC of exon four. The prothrombin cDNA was amplified, digested, and then
CC ligated into RSV-PC at the identical site where the protein C cDNA exons
CC 1-3 had been removed. This construct was then transfected into human
CC 293 cells, from which the chimeric protein can be purified. This

```

CC chimeric protein, can be used as an anticoagulant, to treat disorders where  
 CC protein S is low, some forms of lupus, following stroke or myocardial  
 CC infarction, after venous thrombosis and in disseminated intravascular  
 CC coagulation, adult respiratory distress syndrome, in thromboembolic  
 CC disease or factor V Leiden.

XX  
 SQ Sequence 45 AA;

Query Match 84.9%; Score 169; DB 19; Length 45;  
 Best Local Similarity 93.2%; Pred. No. 1e-20;  
 Matches 41; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 1 ANSFLXLRHSGSLRXRCIXXICDFXXAKXIFZVDVDTLAFWSKH 44  
 |||||  
 DB 1 ANSFLXLRHSGSLRXRCIXXICDFXXAKXIFZVDVDTLAFWSKH 44

RESULT 15

AAV56803  
 ID AAV56803 standard; Protein; 415 AA.

XX  
 AC AAV56803;

XX  
 DT 27-MAR-2000 (first entry)

XX  
 DE Truncated human protein C polypeptide.

XX  
 KM Protein C; truncated; thrombotic disorder; vascular disorder; stroke;  
 KM hypercoagulable state; myocardial infarction; unstable angina; sepsis;  
 KM adult respiratory distress syndrome; sickle cell anemia; human.

XX  
 OS Homo sapiens.

XX  
 PN W09963070-A1.

XX  
 PD 09-DEC-1999.

XX  
 PF 01-JUN-1999; 99WO-US11969.

XX  
 PR 01-JUN-1998; 98US-0087585.

XX  
 PA (ELIL ) LILLY & CO ELI.

XX  
 PI Huang L, Riggin RM;

XX  
 DR WPI: 2000-086975/07.

XX  
 DR N-PSDB: AA246750.

XX  
 PT Novel polypeptide useful for treating thrombotic and vascular diseases  
 PT and hypercoagulation, e.g. stroke

XX  
 PS Claim 2; Page 22-23; 23pp; English.

XX  
 CC This represents a human protein C polypeptide having a light chain and  
 CC a truncated heavy chain. The protein can be produced by standard  
 CC recombinant methodologies. The truncated protein C is used to treat a  
 CC wide range of thrombotic or vascular disorders or hypercoagulable states,  
 CC e.g. stroke; myocardial infarction; unstable angina; sepsis; adult  
 CC respiratory distress syndrome; sickle cell anemia etc. The truncated  
 CC protein C retains the activity of full-length protein C but does not  
 CC undergo C-terminal cleavage, of the heavy chain, during activation.

XX  
 SQ Sequence 415 AA;

Query Match 84.9%; Score 169; DB 21; Length 415;  
 Best Local Similarity 72.7%; Pred. No. 1.1e-19;  
 Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;

OY 1 ANSFLXLRHSGSLRXRCIXXICDFXXAKXIFZVDVDTLAFWSKH 44  
 |||||  
 DB 1 ANSFLXLRHSGSLRXRCIXXICDFXXAKXIFZVDVDTLAFWSKH 44

Search completed: May 13, 2003, 14:03:16  
 Job time : 36 secs

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OW protein - protein search, using sw model

Run on: May 13, 2003, 14:02:03 : Search time 18 seconds  
(without alignments)  
234.995 Million cell updates/sec

Title: SEQ1-EDITED  
Perfect score: 199  
Sequence: 1 ANSFLXLRIGSLXRCIXX.....XXAKXIFZVDVDTLAFWSKH 44

Scoring table: BLOSUM62  
Gapop 10.0, Gapext 0.5

Searched: 283224 seqs, 96134422 residues

Total number of hits satisfying chosen parameters: 283224

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :  
1: PIR\_73:\*  
2: PIR1:\*  
3: PIR3:\*  
4: PIR4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	169	84.9	461	1 KXHU	protein C (activat
2	140	70.4	461	1 JY0210	protein C (activat
3	139	69.8	461	1 S18994	protein C (activat
4	122	61.3	456	1 KXBO	protein C (activat
5	112	56.3	482	1 EXRT	coagulation factor
6	108	54.3	492	1 EXBO	coagulation factor
7	107	53.8	488	1 EXHU	coagulation factor
8	101	50.8	443	2 I65932	coagulation factor
9	98	49.2	466	1 KXHU	coagulation factor
10	84	42.2	407	1 KXBO	coagulation factor
11	83.5	42.0	617	2 S10511	thrombin (EC 3.4.2
12	83.5	42.0	618	2 A35827	thrombin (EC 3.4.2
13	81	40.7	622	1 TBHU	thrombin (EC 3.4.2
14	80	40.2	475	1 EXCH	coagulation factor
15	79	39.7	642	2 S53434	plasma protein S p
16	79	39.7	652	1 KXHU	plasma protein S p
17	78	39.2	476	1 A30351	coagulation factor
18	78	39.2	459	2 J00419	coagulation factor
19	78	39.2	466	2 S38819	plasma protein S -
20	77	38.7	675	1 KXBO	plasma protein S p
21	75	37.7	675	1 KXRTS	coagulation factor
22	74	37.2	461	1 KXHU	plasma protein S p
23	72	36.2	461	1 KXBO	plasma protein S p
24	71	35.7	416	1 KXBO	coagulation factor
25	69	34.7	625	1 TBBO	thrombin (EC 3.4.2
26	68	34.2	675	1 KXMS	plasma protein S p
27	66.5	33.4	396	1 KXBO	plasma protein Z -
28	62.5	31.4	422	1 KXHU	plasma protein Z p
29	59	29.6	673	2 A48089	growth arrest-spec

30	58	29.1	674	2	155476	growth potentialin
31	57	28.6	678	2	B48089	growth arrest-spec
32	56.5	28.4	594	2	D84859	probable MAP kinas
33	54.5	27.4	603	2	C96575	probable MAP kinas
34	52.5	26.4	576	2	G96763	probable MAP kinas
35	52	26.1	606	2	T40556	hypothetical prote
36	49	24.6	730	2	G64062	primosomal replica
37	49	24.6	1363	2	I58375	protein-tyrosine k
38	49	24.6	1684	2	T02367	hypothetical prote
39	49	24.6	1694	2	A83512	hypothetical prote
40	48.5	24.4	510	2	B82918	ammonium transport
41	48	24.1	558	1	S27994	alcohol dehydrogen
42	48	24.1	1235	2	D32433	V56 expression sit
43	48	24.1	1298	2	A48999	protein-tyrosine k
44	47.5	23.9	323	2	T25948	hypothetical prote
45	47	23.6	182	2	JC1189	tyrosine kinase re

## ALIGNMENTS

## RESULT 1

KXHU  
protein C (activated) (EC 3.4.21.69) precursor - human  
N:Alternate names: autoprothrombin IIA; plasma protein C  
C:Species: Homo sapiens (man)  
C>Date: 17-Mar-1987 #sequence\_revision 17-Mar-1987 #text\_change 16-Jul-1999  
C:Accession: A22331; A25426; A21781; A23789; A00927  
R:Foster, D.C.; Yoshitake, S.; Davie, E.W.  
Proc. Natl. Acad. Sci. U.S.A. 82, 4673-4677, 1985  
A>Title: The nucleotide sequence of the gene for human protein C.  
A:Reference number: A22331; MUID:85270390; PMID:2991887  
A:Accession: A22331  
A:Molecule type: DNA  
A:Residues: 1-461 <POS1>  
A:Cross-references: GB:M11228; NID:g190333; PIDN:AAA60166.1; PID:g190334  
R:Plutsky, J.; Hoskins, J.A.; Long, G.L.; Crabtree, G.R.  
Proc. Natl. Acad. Sci. U.S.A. 83, 546-550, 1986  
A>Title: Evolution and organization of the human protein C gene.  
A:Reference number: A25426; MUID:86120978; PMID:3511471  
A:Accession: A25426  
A:Molecule type: DNA  
A:Residues: 1-445, 'L', 446-461 <PLU>  
A:Cross-references: GB:M12712; NID:g190330; PIDN:AAA60165.1; PID:g190332  
R:Foster, D.; Davie, E.W.  
Proc. Natl. Acad. Sci. U.S.A. 81, 4766-4770, 1984  
A>Title: Characterization of a cDNA coding for human protein C.  
A:Reference number: A21781; MUID:84272714; PMID:6589623  
A:Accession: A21781  
A:Molecule type: mRNA  
A:Residues: 'O', 107-461 <FOS2>  
A:Cross-references: GB:X02059; NID:g190322; PIDN:AAA60164.1; PID:g190323  
R:Beckman, R.J.; Schmidt, R.J.; Santet, R.F.; Plutsky, J.; Crabtree, G.R.; Long, G.  
Nucleic Acids Res. 13, 5233-5247, 1985  
A>Title: The structure and evolution of a 461 amino acid human protein C precursor an  
A:Reference number: A23789; MUID:85269639; PMID:2991859  
A:Accession: A23789  
A:Molecule type: mRNA  
A:Residues: 1-461 <BEC>  
A:Cross-references: GB:X02750; NID:g35689; PIDN:CAA6528.1; PID:g763120  
R:Miller, J.P.; Broze Jr., G.J.  
J. Biol. Chem. 265, 11397-11404, 1990  
A>Title: Beta protein C is not glycosylated at asparagine 329. The rate of translatio  
A:Contents: annotation; carbohydrate binding sites; activation peptide  
A>Note: the alpha form of protein C is glycosylated at Asn-329, and the beta form is  
J.Harris, R.J.; Ling, V.T.; Spellman, M.W.  
J. Biol. Chem. 267, 5102-5107, 1992  
A>Title: O-linked fucose is present in the first epidermal growth factor domain of t  
A:Reference number: A44606; MUID:92184750; PMID:1544894  
A:Contents: annotation; beta-hydroxyaspartic acid  
A:Comment: protein C is the zymogen of the vitamin K-dependent serine proteinase that  
ivation of factor Va is strongly enhanced by complexing with protein S. Protein C als

C:Comment: Protein C is synthesized in the liver as a single chain precursor, which is cleaved into two chains, which cleaves a dodecapeptide from the amino end of the heavy chain; this reaction, C:Genetics:

A:Gene: GDB:PROC

A:Cross-references: GDB:120317; OMIM:176860

A:Map position: 2q13-2q21

A:Introns: 24/1; 79/3; 88/1; 134/1; 179/1; 226/3; 266/1

C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology

C:Keywords: anticoagulant; beta-hydroxyaspartic acid; blood coagulation; calcium binding

F:1-32/Domain: signal sequence #status predicted <SIG>

F:27-86/Domain: Gla domain homology <Gla>

F:33-42/Domain: propeptide #status predicted <PRO>

F:43-197/Product: protein C light chain #status predicted <LCH>

F:92-131/Domain: EGF homology <EG1>

F:140-175/Domain: EGF homology <EG2>

F:200-461/Product: protein C heavy chain #status predicted <HCH>

F:212-445/Domain: trypsin homology <TRY>

F:48-49-56-57-60-61-62-67-70-76/Modified site: gamma-carboxyglutamic acid (Glu) #status exp

F:59-64-92-105-101-120-122-131-140-151-147-160-162-175-183-319-238-254-373-387-398-426/D

F:106-111/Disulfide bonds: #status predicted

F:110/Binding site: carbonyl (Thr) (covalent) #status absent

F:113/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status experimental

F:139-290-355/Binding site: carbonyl (Asn) (covalent) #status experimental

F:211-212/Cleavage site: Arg-Leu (thrombin) #status experimental

F:253-299-402/Active site: His, Asp, Ser #status predicted

F:371/Binding site: carbonyl (Asn) (covalent) (partial) #status atypical

Query Match 84.9%; Score 169; DB 1; Length 461;

Best Local Similarity 72.7%; Pred. No. 1.4e-19;

Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSGLRXCIXICDFXXAKXIFDVDDTLAFWSKH 44

DB 43 ANSFLXLRHGSGLRXCIXICDFXXAKXIFDVDDTLAFWSKH 86

RESULT 2

JX0210

protein C (activated) (EC 3.4.21.69) precursor - mouse

N:Alternate names: vitamin K-dependent serine protease

C:Species: Mus musculus (house mouse)

C:Date: 10-Sep-1999 #sequence-revision 10-Sep-1999 #text-change 16-Jun-2000

C:Accession: JX0210

R:Tada, N.; Sato, M.; Tsujimura, A.; Iwase, R.; Hashimoto-Gotoh, T.

J. Biochem. 111, 491-495, 1992

A:Title: Isolation and characterization of a mouse protein C cDNA.

A:Reference number: JX0210; MUID:92316897; PMID:1618739

A:Accession: JX0210

A:Molecule type: mRNA

A:Residues: 1-461 <TAD>

A:Cross-references: GB:DI0445; NID:9220385; PIDN:BA01235.1; PID:9220386

A:Experimental source: liver

C:Comment: Protein C is the zymogen of the vitamin K-dependent serine protease that re

S:

C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homology

C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglut

F:1-33/Domain: signal sequence #status predicted <SIG>

F:27-85/Domain: Gla domain homology <Gla>

F:34-41/Domain: propeptide #status predicted <PRO>

F:42-196-199-461/Product: protein C #status predicted <PRC>

F:91-130/Domain: light chain #status predicted <PCL>

F:139-174/Domain: EGF homology <EG1>

F:199-461/Domain: EGF homology <EG2>

F:199-461/Domain: heavy chain #status predicted <PCH>

F:199-211/Domain: activation peptide #status predicted <ACT>

F:212-445/Domain: trypsin homology <TRY>

F:212-445/Domain: trypsin homology <TRY>

F:47-48-55-57-60-61-66-67-70-76/Modified site: gamma-carboxyglutamic acid (Glu) #status

F:112/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status predicted

F:121-130-139-150-146-159-161-174-182-319-238-254-373-387-398-426/Disulfide bonds: #stat

F:214-230-355/Binding site: carbonyl (Asn) (covalent) #status predicted

F:253-299-402/Active site: His, Asp, Ser #status predicted

Query Match 70.4%; Score 140; DB 1; Length 461;

Best Local Similarity 59.1%; Pred. No. 7.7e-15;

Matches 26; Conservative 7; Mismatches 11; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSGLRXCIXICDFXXAKXIFDVDDTLAFWSKH 44

DB 42 ANSFLXLRHGSGLRXCIXICDFXXAKXIFDVDDTLAFWSKH 85

RESULT 3

S18994

protein C (activated) (EC 3.4.21.69) precursor - rat

C:Species: Rattus norvegicus (Norway rat)

C:Date: 10-Sep-1999 #sequence-revision 10-Sep-1999 #text-change 29-Oct-1999

C:Accession: S18994; S24312

R:Okafuji, T.; Maekawa, K.; Nawa, K.; Marumoto, Y.

submitted to the EMBL Data Library, February 1992

A:Description: The cDNA cloning and mRNA expression of rat protein C.

A:Reference number: S18994

A:Accession: S18994

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-461 <OKA>

A:Cross-references: EMBL:X64336; NID:956962; PIDN:CAA45617.1; PID:956963

R:Okafuji, T.; Maekawa, K.; Nawa, K.; Marumoto, Y.

Biochim. Biophys. Acta 1131, 329-332, 1992

A:Title: The cDNA cloning and mRNA expression of rat protein C.

A:Reference number: S24312; MUID:92325550; PMID:1627650

A:Accession: S24312

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-461 <OKA2>

A:Cross-references: EMBL:X64336; NID:956962; PIDN:CAA45617.1; PID:956963

C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homol

C:Keywords: beta-hydroxyaspartic acid; glycoprotein; hydrolase; serine protease

F:1-32/Domain: signal sequence #status predicted <SIG>

F:27-85/Domain: Gla domain homology <Gla>

F:33-42/Domain: propeptide #status predicted <PRO>

F:43-461/Product: protein C #status predicted <PRC>

F:91-130/Domain: EGF homology <EG1>

F:139-174/Domain: EGF homology <EG2>

F:213-445/Domain: trypsin homology <TRY>

F:47-48-55-57-60-61-66-67-70-76/Modified site: gamma-carboxyglutamic acid (Glu) #stat

F:112/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status predicted

F:121-130-139-150-146-159-161-174-182-320-239-255-373-387-398-426/Disulfide bonds: #s

F:215-291-355/Binding site: carbonyl (Asn) (covalent) #status predicted

F:254-300-402/Active site: His, Asp, Ser #status predicted

Query Match 69.8%; Score 139; DB 1; Length 461;

Best Local Similarity 59.1%; Pred. No. 1.1e-14;

Matches 26; Conservative 7; Mismatches 11; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSGLRXCIXICDFXXAKXIFDVDDTLAFWSKH 44

DB 42 ANSFLXLRHGSGLRXCIXICDFXXAKXIFDVDDTLAFWSKH 85

RESULT 4

KX80

protein C (activated) (EC 3.4.21.69) precursor - bovine (fragment)

N:Alternate names: autoproteolytic protein C; plasma protein C

C:Species: Bos primigenius taurus (cattle)

C:Date: 30-Nov-1980 #sequence-revision 17-Mar-1987 #text-change 16-Jul-1999

C:Accession: A26250; A18385; A18386; A00928

R:Long, G.L.; Balagayle, R.M.; MacGillivray, R.T.A.

Proc. Natl. Acad. Sci. U.S.A. 81, 5653-5656, 1984

A:Title: Cloning and sequence of liver cDNA coding for bovine protein C.

A:Reference number: A26250; MUID:85014826; PMID:6091100

A:Accession: A26250

A:Molecule type: mRNA

A:Residues: 1-456 <LON>

R:Feinlund, P.; Stenflo, J.





A:Cross-references: GB:M22613; NID:g180335; PIDN:AAA51984.1; PID:g180336  
R:Funf, M.R.; Hay, C.W.; MacGillivray, R.T.A.  
A:Title: Characterization of an almost full-length cDNA coding for human blood coagulation  
proc. Natl. Acad. Sci. U.S.A. 82, 3591-3595, 1985  
A:Reference number: A22208; MUID:85216545; PMID:2582420  
A:Accession: A22208  
A:Molecule type: mRNA  
A:Residues: 13-441, 'S', 443-488 <FUN>  
A:Cross-references: GB:K03194; NID:g182840; PIDN:AAA52490.1; PID:g182841  
R:Leyley, S.P.; Chung, D.W.; Kistiel, W.; Kurechi, K.; Davle, E.W.  
A:Title: Characterization of a cDNA coding for human factor X.  
A:Reference number: A21284; MUID:84222026; PMID:6587384  
A:Accession: A21284  
A:Molecule type: mRNA  
A:Residues: 13-284, 'E', 289-488 <LE2>  
A:Cross-references: GB:K01886  
R:McMullen, B.A.; Fujikawa, K.; Kistiel, W.; Sasagawa, T.; Howard, W.N.; Kwa, E.Y.; Weiss  
Biochemistry 22, 2875-2884, 1983  
A:Title: Complete amino acid sequence of the light chain of human blood coagulation factor  
X.  
A:Reference number: A20362; MUID:85257207; PMID:6871167  
A:Accession: A20362  
A:Molecule type: Protein  
A:Residues: 41-179 <MCN>  
R:Inoue, K.; Morita, T.  
Eur. J. Biochem. 218, 153-163, 1993  
A:Title: Identification of O-linked oligosaccharide chains in the activation peptides of  
A:Reference number: S39414; MUID:94062825; PMID:8234461  
A:Accession: S39415  
A:Molecule type: protein  
A:Residues: 183-234 <INO>  
A:Note: glycosylation sites  
A:Note: Identification and characterization of beta-hydroxyaspartic acid  
R:Jagadeeswaran, P.; Reddy, S.V.; Rao, K.J.; Hamsabhusanam, K.; Lyman, G.  
Gene 84, 517-519, 1989  
A:Title: Cloning and characterization of the 5' end (exon 1) of the gene encoding human  
A:Reference number: I54051; MUID:90128299; PMID:2612918  
A:Accession: I54051  
A:Status: translation not shown; translated from GB/EMBL/DBJ  
A:Molecule type: DNA  
A:Residues: 1-23 <RES>  
A:Cross-references: GB:M3297; NID:g183860; PIDN:AAA52636.1; PID:g553330  
R:Padmanabhan, K.; Padmanabhan, K.P.; Tullinsky, A.; Park, C.H.; Bode, W.; Huber, R.; Blumberg  
J. Mol. Biol. 232, 947-966, 1993  
A:Title: Structure of human des(1-45) factor Xa at 2.2 angstroms resolution.  
A:Reference number: A49458; MUID:93360277; PMID:8355279  
A:Contents: annotation: X-ray crystallography, 2.2 angstroms  
A:Comment: The two chains held together by one disulfide bond are formed from a single-c  
A:Comment: The activation peptide is cleaved by factor IXa (in the intrinsic pathway) or  
A:Genetics:  
A:Gene: GDB:F10  
A:Cross-references: GDB:119890; OMIM:227600  
A:Map position: 13q34-13q34  
A:Introns: 24/1; 77/3; 86/1; 124/1; 150/3; 249/3; 289/1  
A:Note: deficiency of this factor causes Stuart disease  
A:Function:  
A:Description: catalyzes the proteolytic activation of prothrombin to thrombin in the pr  
A:Pathway: blood coagulation  
C:Superfamily: blood coagulation  
C:Keywords: beta-hydroxyaspartic acid, blood coagulation; calcium binding; carboxylat  
E:1-23/Domain: signal sequence #status predicted <SIG>  
E:24-40/Domain: propeptide #status predicted <PRO>  
E:25-84/Domain: Gla domain homology <GLA>  
E:41-179/Product: coagulation factor X light chain #status experimental <LCN>  
E:90-121/Domain: EGF homology <EGF1>  
E:129-164/Domain: EGF homology <EG2>  
E:183-488/Product: coagulation factor X heavy chain #status experimental <HCN>  
E:183-234/Domain: activation peptide #status experimental <APT>  
E:235-488/Product: coagulation factor Xa heavy chain #status experimental <ACT>  
E:235-462/Domain: trypsin homology <TRY>  
E:46-47, 54, 56, 59, 60, 65, 66, 69, 72, 79/Modified site: gamma-carboxyglutamic acid (Glu) #sta  
E:57-62/Disulfide bonds: #status predicted  
E:90-101, 95-110, 112-121, 129-140, 130-149, 151-164, 172-342, 241-246, 261-277, 390-404, 415-443, 4

F:103/Modified site: erythro-beta-hydroxypaspatic acid (Asp) #status experimental  
F:199,231/Binding site: carbohydrate (Thr) (covalent) #status experimental  
F:221,231/Binding site: carbohydrate (Asn) (covalent) #status experimental  
F:234-235/Cleavage site: Arg-Tle (coagulation factor IXa, coagulation factor VIIa  
F:276,322,419/Active site: His, Asp, Ser #status experimental

Query Match	53.8%	Score 107;	DB 1;	Length 488;
Best Local Similarity	40.9%;	Pred. No. 2e-09;		
Matches	18;	Conservative	9;	Mismatches 17;
			Indels	0;
			Gaps	0

QY 1 ANSFLXXLRHGSGLARXCIXXICDFXAAXKIFZDVDDTLAFWSKH 44  
||| : | | | : | : | : ||| :  
Db 41 ANSFLLEMKKGHLIERECMEETCSYEAREVFEDSDKTNEFWNKY 84

RESULT 8  
TAS033

coagulation factor VII - rabbit

C:\species: Oryctolagus cuniculus (domestic rabbit)  
C:\Date: 04-Sep-1997 #sequence revision 04-Sep-1997 #text change 12-Feb-1998

C;Accession: I46932

Thromb. Res. 69, 231-238, 1993

A/Title: Complete nucleotide sequence of the cDNA encoding the human TACC3 with 330000 bp

A;Accession: I46932

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Residues: 1-443 <BR>

A; Cross-references: GB:538300; NID:g266294; PID:g266293  
 C: Superfamily: coagulation factor X; EGF homology: G1A

F; 24-83/Domain: Gla domain homology <GLA>

F:130-166/Domain: EGF homology <EG2>

F;192-425/Domain: trypsin homology <TRY>

Query Match	50.88; Score 101; DB 2;
-------------	-------------------------

Best Local Similarity 46.38; Pred. No. 1.8e-08;  
Matches 19: Conservative 5; Mismatches 17; Indels 0  
Cross 0

1 ANSE EAAENHSEAKCALIAICDF AAAAATF ZDVDDI LAEW 41

Db 40 ANSFLLELRPGSLERECKEELCSFEAREVFQSTERTKQFW 80

## RESULT 9

coagulation factor VIIa (EC 3.4.21.21) precursor [validated] - human

C:Species: Homo sapiens (man)  
C:Date: 10-Mar-1080 #seconds  
C:Date: 10-Mar-1000 #seconds  
C:Date: 00-Dec-2000

C;Accession: A28322; A23819; A31186; B31186; S63524

R;U Hata, P.O.; Grant, F.J.; Haldeman, B.A.; Gray, C.L.; Insley, M.Y.; Hage  
Proc Natl Acad Sci U S A 84 5158-5162 1987

A:Title: Nucleotide sequence of the gene coding for human factor VII, a vit

A:Accession: A28322  
n/reference number: A20322, MOLD:0/200340, PMID:303/33/

A;Molecule type: DNA

A;Cross-references: GB:J02933; NID:q180333; PIDN:AAA51983.1; PID:q180334

R;Hagen, F.S.; Gray, C.L.; O'Hara, P.; Grant, F.J.; Saari, G.C.; Woodbury, D.; Na+1 Acad Sci U S A 82, 2415-2416, 1985

A;Title: Characterization of a cDNA coding for human factor VII.

A:Accession: A23819  
A:Reference Number: A23819; MOLD:86205965; PMID:3486420

A; molecule type: mRNA

A:Cross-references: GB:M13232: NID:q182799: PIDN:AAA88040.1: PTD:q182801

R;Thlm, L.; Bjoern, S.; Christensen, M.; Nicolaisen, E.M.; Lund-Hansen, T.;  
Elschert, H. 27 7705-7707 1000

A;Title: Amino acid sequence and posttranslational modifications of human f

A: Accession: A31186  
A: Reference number: AY0539; MUID:89088153; PMID:3264/25

A;Molecule type: protein



A:Residues: 61-212 <TH1>  
A:Accession: B31166  
A:Molecule type: protein  
A:Residues: 213-466 <TH2>  
R:Björner, S.; Foster, D.C.; Thim, L.; Wülfberg, F.C.; Christensen, M.; Komiyama, Y.; Pedersen, J. Biol. Chem. 266, 11051-11057, 1991  
A:Title: Human plasma and recombinant factor VII. Characterization of O-glycosylations and contents: annotation; carbohydrate binding sites  
R:Petersen, E.; Petersen, L.C.  
Eur. J. Biochem. 234, 293-300, 1995  
A:Title: Structurally and functionally distinct Ca(2+) binding sites in the gamma-carboxyglutamate domain of factor VII  
A:Reference number: S63524; MUID:96096752; PMID:8529655  
A:Accession: S63524  
A:Molecule type: protein  
A:Residues: 61-65;99-103;105-109;213-308-312 <PER>  
C:Genetics:  
A:Gene: GDB:F7  
A:Cross-references: GDB:119897; OMIM:227500  
A:Map position: 13q34-13q34  
A:Introns: 22/1; 44/1; 97/3; 106/1; 144/1; 191/1; 227/3; 269/1  
C:Function:  
A:Description: catalyzes the proteolytic activation of coagulation factor X in the presence of calcium and tissue factor  
A:Pathway: blood coagulation extrinsic pathway  
C:Keywords: beta-hydroxyaspartic acid; blood coagulation; Gla domain homology; trypsin homology  
F:1-20/Domain: signal sequence #status predicted <SIG>  
F:21-60/Domain: propeptide #status predicted <PRO>  
F:61-104/Domain: Gla domain homology <Gla>  
F:61-212/Product: coagulation factor VIIa light chain #status experimental <MA1>  
F:110-141/Domain: EGF homology <EG1>  
F:151-187/Domain: EGF homology <EG2>  
F:213-466/Product: coagulation factor VIIa heavy chain #status experimental <MA2>  
F:213-447/Domain: trypsin homology <TRY>  
F:66-67;74;76;79;80;85;86;89;95/Modified site: gamma-carboxyglutaminc acid (Glu) #status experimental  
F:77-82;110-121;115-130;132-141;151-162;158-172;174-187;195-322;219-224;238-254;370-389;  
F:112,120/Binding site: carbohydrate (Ser) (covalent) #status experimental  
F:123/Modified site: erythro-beta-hydroxyaspartic acid (Asp) #status absent  
F:205;382/Binding site: carbohydrate (Asn) (covalent) #status experimental  
F:212-213/Cleavage site: Arg-Ile (coagulation factor XIla) #status experimental  
F:253;302;404/Active site: His, Asp, Ser #status predicted  
F:350-351/Cleavage site: Arg-Gly (coagulation factor XIa) #status predicted

Query Match 49.2%; Score 98; DB 1; Length 466;  
Best Local Similarity 48.8%; Pred. No. 5.8e+08;  
Matches 20; Conservative 4; Mismatches 17; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSILKRCIXICDFXXAKXIFZDVDDTLAFW 41  
II:II II III I I I I I I I I I I I I I I I I I I  
Db 61 ANAFLEELRPGSLERECKEEOCSFEAREIFPKDARTKLFW 101  
RESUTJ 10  
RKBO7  
coagulation factor VIIa (EC 3.4.21.21) - bovine  
C:Species: Bos primigenius taurus (cattle)  
C:Date: 21-May-1990 sequence\_revision 23-Mar-1995 #text\_change 16-Jul-1999  
C:Accession: A31979; C20774  
R:Takeya, H.; Kawabata, S.; Nakagawa, K.; Yamamichi, Y.; Miyata, T.; Iwanaga, S.  
J. Biol. Chem. 263, 14866-14877, 1988  
A:Title: Bovine factor VII. Its purification and complete amino acid sequence.  
A:Reference number: A31979; MUID:89008362; PMID:3049594  
A:Accession: A31979  
A:Molecule type: protein  
A:Residues: 1-407 <TKA>  
R:McMullen, B.A.; Fujikawa, K.; Kisiel, W.  
Biochem. Biophys. Res. Commun. 115, 8-14, 1983  
A:Title: The occurrence of beta-hydroxyaspartic acid in the vitamin K-dependent blood coagulation factors  
A:Reference number: A20274; MUID:83308813; PMID:6688526  
A:Accession: C20274

A:Note: the residue designated 'X' was determined to be hydroxyaspartic acid  
R.Hase, S.; Kanabata, S.; Nishimura, H.; Takaya, H.; Sueyoshi, T.; Miyata, T.; Iwanag  
J. Biochem. 104, 867-868, 1988  
A:Title: A new trisaccharide sugar chain linked to a serine residue in bovine blood c  
A:Reference number: A44556; MUID:89213999; PMID:3149637  
A:Contents: annotation  
A:Note: structure and location of covalently bound carbohydrate  
C:Function:  
A:Description: catalyzes the proteolytic activation of coagulation factor X in the pr  
gulation factor IX in the presence of calcium and tissue factor  
A:Pathway: blood coagulation extrinsic pathway  
C:Superfamily: coagulation factor X; EGF homology; Gla domain homology; trypsin homol  
C:Keywords: beta-hydroxyaspartic acid; blood coagulation; calcium binding; carboxyglu  
F:1-152/Product: coagulation factor VIIa light chain #status experimental <MA1>  
F:1-44/Domain: Gla domain homology (fragment) <GLA>  
F:50-81/Domain: EGF homology <EG1>  
F:91-127/Domain: EGF homology <EG2>  
F:153-407/Product: coagulation factor VIIa heavy chain #status experimental <MA2>  
F:153-387/Domain: trypsin homology <TRY>  
F:6-7,14,16,19,20,25,26,29,34,35/Modified site: gamma-carboxyglutamic acid (Glu) #Sta  
F:17-22,50-61,55-70,72-81,91-102,98-112,114-127,135-262,159-164,178-194,310-329,340-3  
F:52/Binding site: carboxylate (Ser) (covalent) #status experimental  
F:152/Modified site: erythro-beta-hydroxyaspartic acid (Asp) (partial) #status experim  
F:145-203/Binding site: carboxylate (Asn) (covalent) #status experimental  
F:152-153/Binding site: Arg-Tile (coagulation factor X) #status experimental  
F:193-242,344/Active site: His, Asp, Ser #status predicted  
F:290-291/Cleavage site: Arg-Gly (coagulation factor Xa) #status experimental

Query Match 42.2%; Score 84; DB 1; Length 407;  
Best Local Similarity 43.9%; Pred. No. 9, 86-06;  
Matches 18; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 ANSFLXLRHGSGLKRXCIXXICDFYXKXIFZDVDDTLAFW 41  
||| | ||| | : | | | | : | | |  
Db 1 ANGFLLELLPGSLRECRCELCSFEAEHIFRNEERTQFW 41

RESULT 11  
S10511  
thrombin (EC 3.4.21.5) precursor - rat  
C:Species: Rattus norvegicus (Norway rat)  
C:Date: 07-May-1993 #sequence revision 07-May-1993 #text\_change 03-May-2002  
C:Accession: S10511; A60576; B42696  
R:Diwanich, M.; Monard, D.  
Nucleic Acids Res. 18, 4251, 1990  
A:Title: cDNA sequence of rat prothrombin.  
A:Reference number: S10511; MUID:90332426; PMID:2377469  
A:Accession: S10511  
A:Molecule type: mRNA  
A:Residues: 1-617 <DHI>  
A:Cross-references: EMBL:X52835; NID:956969; PIDN:CA437017.1; PID:956970  
R:Henriksson, K.P.; Jazin, E.E.; Greenwood, J.A.; Dickerman, H.W.  
Endocrinology 126, 167-175, 1990  
A:Title: Prothrombin levels are increased in the estrogen-treated immature rat uterus  
A:Reference number: A60576; MUID:90091942; PMID:2293980  
A:Accession: A60576  
A:Molecule type: protein  
A:Residues: 44-58 <HEN>  
R:Banfield, D.K.; MacGillivray, R.T.A.  
Proc. Natl. Acad. Sci. U.S.A. 89, 2779-2783, 1992  
A:Title: Partial characterization of vertebrate prothrombin cDNAs: amplification and  
A:Reference number: A42696; MUID:92212913; PMID:1557383  
A:Accession: B42696  
A:Status: preliminary  
A:Molecule type: mRNA  
A:Residues: 383-617, 'E' <BAN>  
A:Cross-references: GB:M81397  
C:Superfamily: thrombin; Gla domain homology; kinase homology; trypsin homology  
C:Keywords: blood coagulation; calcium binding; carboxyglutamic acid; glycoprotein; h  
F:1-24/Domain: signal sequence #status predicted <SIG>  
F:25-43/Domain: propeptide #status predicted <PRO>  
F:28-88/Domain: Gla domain homology <GLA>







GenCore version 5.1.4-p5-4578  
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OM protein - protein search, using sw model

Run on: May 13, 2003, 14:01:08 ; Search time 11 seconds  
(without alignments)  
165.905 Million cell updates/sec

Title: SEQ1-EDITED  
Perfect score: 199  
Sequence: 1 ANSFLXLRHGSLSLRXCIX.....XXAKXIFZVDDTLAFWSKH 44

Scoring table: BLOSUM62  
Gapop 10.0 , Gapext 0.5

Searched: 112892 seqs, 41476328 residues

Total number of hits satisfying chosen parameters: 112892

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database : SwissProt\_40.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	169	84.9	461	1 PRTC_HUMAN	P04070 homo sapien
2	140	70.4	461	1 PRTC_MOUSE	P33387 mus musculu
3	139	69.8	461	1 PRTC_RAT	P31394 rattus norv
4	138	69.3	458	1 PRTC_RABIT	Q28661 oryctolagus
5	123	61.8	459	1 PRTC_PIG	Q9G1P2 sus scrofa
6	122	61.3	456	1 PRTC_BOVIN	P00743 bos taurus
7	108	54.3	492	1 FA10_BOVIN	P00743 bos taurus
8	107	53.8	488	1 FA10_HUMAN	P00742 homo sapien
9	107	50.8	231	1 TMG3_HUMAN	Q9BZD7 homo sapien
10	101	50.8	444	1 FA7_RABIT	P98139 oryctolagus
11	100	50.3	490	1 FA10_RABIT	O19045 oryctolagus
12	98	49.2	465	1 FA7_HUMAN	P08709 homo sapien
13	86	43.2	218	1 TMG1_HUMAN	O14668 homo sapien
14	84	42.2	226	1 TMG4_HUMAN	Q9BZD6 homo sapien
15	84	42.2	407	1 FA7_BOVIN	P18292 rattus norv
16	83.5	42.0	617	1 THRB_RAT	P19221 mus musculu
17	83.5	42.0	618	1 THRB_MOUSE	P00734 homo sapien
18	81	40.7	622	1 THRB_HUMAN	P81428 tropidichis
19	80	40.2	376	1 FA10_TROCA	P25155 gallus gall
20	80	40.2	475	1 FA10_CHICK	P25155 gallus gall
21	79	39.7	446	1 FA7_MOUSE	P70375 mus musculu
22	79	39.7	649	1 PRTS_MACMU	Q28520 macaca mula
23	79	39.7	649	1 PRTS_HUMAN	P07225 homo sapien
24	78	39.2	452	1 FA9_MOUSE	P19540 canis fami
25	78	39.2	459	1 FA9_MOUSE	P16294 mus musculu
26	78	39.2	456	1 PRTS_RABIT	P98118 oryctolagus
27	77	38.7	675	1 PRTS_BOVIN	P07224 bos taurus
28	74	37.2	461	1 PRTS_RAT	P53813 rattus norv
29	74	37.2	461	1 FA9_HUMAN	P00741 homo sapien
30	71	35.7	416	1 THRB_BOVIN	P00735 bos taurus
31	69	34.7	625	1 PRTS_MOUSE	O08761 mus musculu
32	68	34.2	675	1 PRTS_MOUSE	P00744 bos taurus
33	66.5	33.4	396	1 PRT2_BOVIN	

34	63	31.7	202	1 TMG2_HUMAN	O14669 homo sapien
35	62.5	31.4	400	1 PRTC_HUMAN	P22891 homo sapien
36	50	25.1	501	1 MKC1_CANAL	P43068 canida alb
37	49	24.6	730	1 PRTA_HAEIN	P44647 haemophilus
38	49	24.6	1363	1 VGR1_MOUSE	P35917 mus musculu
39	48	24.1	358	1 ALK1_PSEOL	Q00593 pseudomonas
40	48	24.1	1235	1 CYA4_TRYBB	Q26721 trypanosoma
41	48	24.1	1298	1 VGR3_HUMAN	P35916 homo sapien
42	47	23.6	1343	1 VGR2_RAT	O08775 rattus norv
43	47	23.6	1348	1 VGR2_COTJA	P52583 coturnix co
44	47	23.6	1356	1 VGR2_HUMAN	P35968 homo sapien
45	47	23.6	1367	1 VGR2_MOUSE	P35918 mus musculu

## ALIGNMENTS

RESULT 1  
PRTC\_HUMAN STANDARD: PRT; 461 AA.  
AC P04070: Q16001: Q15190: Q15189;  
DT 01-NOV-1986 (Rel. 03, Created)  
DT 01-NOV-1986 (Rel. 03, Last sequence update)  
DT 15-JUN-2002 (Rel. 41, Last annotation update)  
DE Vitamin-K dependent protein C precursor (EC 3.4.21.69)  
DE (Autoproteolysin IIA) (Anticoagulant protein C) (Blood coagulation factor XIV).  
CN PROC.  
OS Homo sapiens (Human).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.  
OX NCBI\_TaxID=9606;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=85270390; PubMed=2991887;  
RA Foster D.C., Yoshitake S., Davie E.W.;  
RL Proc. Natl. Acad. Sci. U.S.A. 82:4673-4677(1985).  
RN [2]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=85269639; PubMed=2991859;  
RA Beckmann R.J., Schmidt R.J., Santerre R.F., Plutsky J., Crabtree G.R., Long G.L.;  
RL "The structure and evolution of a 461 amino acid human protein C precursor and its messenger RNA, based upon the DNA sequence of cloned human liver cDNAs";  
RN Nucleic Acids Res. 13:5233-5247(1985).  
RN [3]  
RP SEQUENCE FROM N.A.  
RX MEDLINE=86120978; PubMed=3511471;  
RA Plutsky J., Hoskins J.A., Long G.L., Crabtree G.R.;  
RL "Evolution and organization of the human protein C gene";  
RN Proc. Natl. Acad. Sci. U.S.A. 83:346-350(1986).  
RN [4]  
RP SEQUENCE FROM N.A.  
RX Rieder M.J., Cartington D.P., Chung M.-W., Lee K.L., Pool C.L., Yi Q., Nickerson D.A.;  
RL Submitted (JUN-2001) to the EMBL/GenBank/DBJ databases.  
RN [5]  
RP SEQUENCE OF 106-461 FROM N.A.  
RX MEDLINE=84272714; PubMed=6589623;  
RA Foster D.C., Davie E.W.;  
RL "Characterization of a cDNA coding for human protein C";  
RN Proc. Natl. Acad. Sci. U.S.A. 81:4766-4770(1984).  
RN [6]  
RP CARBOHYDRATE-LINKAGE SITE ASN-371.  
RX MEDLINE=90293094; PubMed=1694179;  
RA Mielich J.P., Broze G.J. Jr.;  
RL "Beta protein C is not glycosylated at asparagine 329. The rate of translation may influence the frequency of usage at asparagine-X-cysteine sites";  
RN J. Biol. Chem. 265:11397-11404(1990).  
RN [7]

RP HYDROXYLATION.  
 RX MEDLINE-92184750; PubMed-1544894;  
 RA Harris R.J., Ling V.T., Spellman M.W.;  
 RT "O-linked fucose is present in the first epidermal growth factor  
 RL domain of factor XII but not protein C.";  
 RN J. Biol. Chem. 267:5102-5107(1992).  
 RM (18)  
 RP 3D-STRUCTURE MODELING OF 175-450.  
 RX MEDLINE-94272342; PubMed-8003977;  
 RA Fisher C.L., Greengard J.S., Griffin J.H.;  
 RT "Models of the serine protease domain of the human antithrombotic  
 RL plasma factor activated protein C and its zymogen.";  
 RN Protein Sci. 3:588-599(1994).  
 RM (19)  
 RP X-RAY CRYSTALLOGRAPHY (2.8 ANGSTROMS) OF 84-461.  
 RX MEDLINE-97157472; PubMed-9003757;  
 RA Mather T., Oganessyan V., Hof P., Huber R., Foundling S., Esmen C.,  
 RL Bode W.;  
 RN "The 2.8 A crystal structure of Gla-domainless activated protein C.";  
 RM EMBO J. 15:6822-6831(1996).  
 RL (10)  
 RP REVIEW ON PROC VARIANTS.  
 RX MEDLINE-93190290; PubMed-8446940;  
 RA Reltsma P.H., Poort S.R., Bernardi F., Gandrille S., Long G.L.,  
 RL Sala N., Cooper D.N.;  
 RN "Protein C deficiency: a database of mutations. For the Protein C & S  
 RT Subcommittee of the Scientific and Standardization Committee of the  
 RL International Society on Thrombosis and Haemostasis.";  
 RM Thromb. Haemost. 69:77-84(1993).  
 RL (11)  
 RP VARIANT CYS-444.  
 RX MEDLINE-87204221; PubMed-2437584;  
 RA Romeo G., Hassan H.T., Staempfli S., Roncuzzi L., Cianetti L.,  
 RL Leonard A., Vicente V., Mannucci P.M., Bertina R.M., Peschle C.,  
 RN Cortese R.;  
 RM "Hereditary thrombophilia: identification of nonsense and missense  
 RT mutations in the protein C gene.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 84:2829-2832(1987).  
 RM (12)  
 RP VARIANT TRP-211 (LONDON-1).  
 RX MEDLINE-90098906; PubMed-2602169;  
 RA Grundy C.B., Chitcolle A., Talbot S., Bevan D., Kakkar V.V.,  
 RL Cooper D.N.;  
 RN "Protein C London 1: recurrent mutation at Arg-169 (CGG->TGC) in  
 RT the protein C gene causing thrombosis.";  
 RL Nucleic Acids Res. 17:10513-10513(1989).  
 RM (13)  
 RP VARIANT CYS-272.  
 RX MEDLINE-91329836; PubMed-1868249;  
 RA Reltsma P.H., Poort S.R., Allart C.F., Briet E., Bertina R.M.;  
 RL "The spectrum of genetic defects in a panel of 40 Dutch families with  
 RT symptomatic protein C deficiency type I: heterogeneity and founder  
 RL effects.";  
 RN Blood 78:890-894(1991).  
 RL (14)  
 RP VARIANTS ALA-62 (VERMONT-1) AND MET-76.  
 RX MEDLINE-92190481; PubMed-1347706;  
 RA Bovill E.G., Tomczak J.A., Grant B., Bhushan F., Pillmer E.,  
 RL Raluville I.R., Long G.L.;  
 RN "Protein C Vermont: symptomatic type II protein C deficiency  
 RT associated with two GLA domain mutations.";  
 RL Blood 79:1456-1465(1992).  
 RM (15)  
 RP VARIANT ASP-418 (HONG KONG-2).  
 RX MEDLINE-92305321; PubMed-161081;  
 RA Sugahara Y., Miura O., Yuen P., Aoki N.;  
 RL "Protein C deficiency Hong Kong 1 and 2: hereditary protein C  
 RT deficiency caused by two mutant alleles, a 5-nucleotide deletion and  
 RL a missense mutation.";  
 RN Blood 80:126-133(1992).  
 RM (16)  
 RP VARIANT LEU-289.  
 RX MEDLINE-92380660; PubMed-1511988;

RA Grundy C.B., Chisholm M., Kakkar V.V., Cooper D.N.;  
 RT "A novel homozygous missense mutation in the protein C (PROC) gene  
 RL causing recurrent venous thrombosis.";  
 RN Hum. Genet. 89:683-684(1992).  
 RM (17)  
 RP VARIANTS GLN-220 AND TRP-220.  
 RX MEDLINE-92380661; PubMed-1511989;  
 RA Grundy C.B., Schuilen S., Tengborn L., Kakkar V.V., Cooper D.N.;  
 RT "Two different missense mutations at Arg 178 of the protein C (PROC)  
 RL gene causing recurrent venous thrombosis.";  
 RN Hum. Genet. 89:685-686(1992).  
 RM (18)  
 RP VARIANT GLN-220.  
 RX MEDLINE-93250852; PubMed-1301959;  
 RA Gandrille S., Vidard M., Atsch M., Alhenc-Gelas M., Fischer A.M.,  
 RL Couault-Hellman M., Toulon P., Flessinger J.N., Goossens M.;  
 RN "Two novel mutations responsible for hereditary type I protein C  
 RT deficiency: characterization by denaturing gradient gel  
 RL electrophoresis.";  
 RM Hum. Mutat. 1:491-500(1992).  
 RL (19)  
 RP VARIANT SER-334.  
 RX MEDLINE-92276939; PubMed-1593215;  
 RA Yamamoto K., Matsushita T., Sugitara I., Takamatsu J., Iwasaki E.,  
 RL Wada H., Deguchi K., Shirakawa S., Saito H.;  
 RN "Homozygous protein C deficiency: identification of a novel missense  
 RT mutation that causes impaired secretion of the mutant protein C.";  
 RL J. Lab. Clin. Med. 119:682-689(1992).  
 RM (20)  
 RP VARIANTS TRP-38; CYS-42; HIS-42; GLN-271 AND ASN-294.  
 RX MEDLINE-93313192; PubMed-8324221;  
 RA Gandrille S., Alhenc-Gelas M., Gaussem P., Allard M.-F., Dupuy E.,  
 RL Juhan-Vague I., Atsch M.;  
 RN "Five novel mutations located in exons III and IX of the protein C  
 RT gene in patients presenting with defective protein C anticoagulant  
 RL activity.";  
 RM Blood 82:159-168(1993).  
 RL (21)  
 RP VARIANTS G-14; Q-211; Y-244; Q-253; L-321; C-328; I-385; T-388 AND  
 RV-388.  
 RX MEDLINE-93271391; PubMed-8499565;  
 RA Poort S.R., Pabinger-Fasching I., Mannhalter C., Reltsma P.H.,  
 RL Bertina R.M.;  
 RN "Twelve novel and two recurrent mutations in 14 Austrian families  
 RT with hereditary protein C deficiency.";  
 RL Blood Coagul. Fibrinolysis 4:273-280(1993).  
 RM (22)  
 RP VARIANT TRP-57.  
 RX MEDLINE-93271396; PubMed-8499568;  
 RA Millar D.S., Grundy C.B., Bignell P., Moffat E.H., Martin R.,  
 RL Kakkar V.V., Cooper D.N.;  
 RN "A Gla domain mutation (Arg 15-->Trp) in the protein C (PROC) gene  
 RT causing type 2 protein C deficiency and recurrent venous  
 RL thrombosis.";  
 RM Blood Coagul. Fibrinolysis 4:345-347(1993).  
 RL (23)  
 RP VARIANTS R-145; L-210; W-211; T-243; L-321; M-340 AND Y-426.  
 RX MEDLINE-94122329; PubMed-8292730;  
 RA Tsay W., Greengard J.S., Montgomery R.R., McPherson R.A., Fucci J.C.,  
 RL Koepfer M.A., Coughlin J., Griffin J.H.;  
 RN "Genetic mutations in ten unrelated American patients with  
 RT symptomatic type I protein C deficiency.";  
 RL Blood Coagul. Fibrinolysis 4:791-796(1993).  
 RM (24)  
 RP VARIANT SER-423.  
 RX MEDLINE-94001606; PubMed-8398832;  
 RA Marchetti G., Patrachini P., Gemmati D., Castaman G., Rodeghiero F.,  
 RL Wacey A., Cooper D.N., Tuddenham E.G., Bernardi F.;  
 RN "Symptomatic type II protein C deficiency caused by a missense  
 RT mutation (Gly 381-->Ser) in the substrate-binding pocket.";  
 RL Br. J. Haematol. 84:285-289(1993).  
 RM (25)  
 RP SEQUENCE OF 43-64 FROM N.A., AND VARIANT GLY-57 (YONAGO).













```

FT DISULFID 159 172 BY SIMILARITY.
FT DISULFID 180 318 INTERCHAIN.
FT DISULFID 237 253
FT DISULFID 368 382
FT DISULFID 393 421
FT CARBOHYD 136 136 N-LINKED (GLCNAC. . .)
FT CARBOHYD 289 289 N-LINKED (GLCNAC. . .)
FT CARBOHYD 350 350 N-LINKED (GLCNAC. . .)
FT CARBOHYD 366 366 N-LINKED (GLCNAC. . .)
FT VARIANT 82 82 F -> K.
FT CONFLICT 455. 456 VP -> PV (IN REF. 4).
SQ SEQUENCE 456 AA; 51407 MW; CAAFE833F894C209 CRC64;

Query Match 61.3%; Score 122; DB 1; Length 456;
Best Local Similarity 50.0%; Pred. No. 1.5e-13;
Matches 21; Conservative 9; Mismatches 12; Indels 0; Gaps 0;

QY 1 ANSFLXLRHGSIXRXIXXICDFXKAKXIFZDVDTLAWFS 42
Db 40 ANSFLELRPGNVERECSEFEERELFONTEDTMAFWS 81

RESULT 7
FA10_BOVIN STANDARD; PRT; 492 AA.
ID FA10_BOVIN
AC P00743;
DT 21-JUL-1986 (Rel. 01, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Coagulation factor X precursor (EC 3.4.21.6) (Stuart factor).
GN F10.
OS Bos taurus (Bovine).
OC Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN 11
RX SEQUENCE OF 1-487 FROM N.A.
RA MEDLINE=84247315; PubMed=6330671;
RA Fung M.R., Campbell R.M., McGillivray R.T.A.;
RT "Blood coagulation factor X mRNA encodes a single polypeptide chain
RT containing a prepro leader sequence."
RL Nucleic Acids Res. 12:4461-4492(1984).
RN 12
RX SEQUENCE OF 41-180.
RA MEDLINE=80130563; PubMed=6766735;
RA Enfield D.L., Ericsson L.H., Fujikawa K., Walsh K.A., Neurath H.,
RA Titani K.;
RT "Amino acid sequence of the light chain of bovine factor XI (Stuart
RT factor).";
RL Biochemistry 19:659-667(1980).
RN 13
RX REVISION TO 103.
RA MEDLINE=83308813; PubMed=6688526;
RA McMullen B.A., Fujikawa K., Kistiel W.;
RT "The occurrence of beta-hydroxyaspartic acid in the vitamin
RT K-dependent blood coagulation zymogens."
RL Biochem. Biophys. Res. Commun. 115:8-14(1983).
RN 14
RX SEQUENCE OF 183-492, CARBOHYDRATE-LINKAGE SITES, AND DISULFIDE BONDS.
RA MEDLINE=76053069; PubMed=1059093;
RA Titani K., Fujikawa K., Enfield D.L., Ericsson L.H., Walsh K.A.,
RA Neurath H.;
RT "Bovine factor XI (Stuart factor): amino-acid sequence of heavy
RT chain.";
RL Proc. Natl. Acad. Sci. U.S.A. 72:3082-3086(1975).
RN 15
RX SEQUENCE OF 183-233, AND CARBOHYDRATE-LINKAGE SITES.
RA MEDLINE=94062825; PubMed=8243461;
RA Inoue K., Morita T.;
RT "Identification of O-linked oligosaccharide chains in the activation
RT peptides of blood coagulation factor X. The role of the carbohydrate
RT moieties in the activation of factor X.";

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RL Eur. J. Biochem. 218:153-163(1993).
RN [6]
RX ACTIVE SITE.
RA MEDLINE=73053314; PubMed=4264286;
RA Titani K., Hermanson M.A., Fujikawa K., Ericsson L.H., Walsh K.A.,
RA Neurath H., Davie E.W.;
RT "Bovine factor X 1a (activated Stuart factor). Evidence of homology
RT with mammalian serine proteases.";
RN Biochemistry 11:4899-4903(1972).
RN [7]
RX PROCESSING.
RA MEDLINE=76053121; PubMed=1059122;
RA Fujikawa K., Titani K., Davie E.W.;
RT "Activation of bovine factor X (Stuart factor): conversion of factor
RT Xa-alpha to factor Xa-beta.";
RN Proc. Natl. Acad. Sci. U.S.A. 72:3359-3363(1975).
RN [8]
RX CALCIUM-BINDING DATA.
RA MEDLINE=84185716; PubMed=6546930;
RA Sugo T., Bjoerk I., Holmgren A., Stenflo J.;
RT "Calcium-binding properties of bovine factor X lacking the gamma-
RT carboxyglutamic acid-containing region.";
RN J. Biol. Chem. 259:5705-5710(1984).
RN [9]
RX SULFATION.
RA MEDLINE=86140210; PubMed=3949800;
RA Morita T., Jackson C.M.;
RT "Localization of the structural difference between bovine blood
RT coagulation factors XI and X2 to tyrosine 18 in the activation
RT peptide.";
RN J. Biol. Chem. 261:4008-4014(1986).
RN [10]
RX STRUCTURE BY NMR OF 85-126.
RA MEDLINE=91084483; PubMed=2261466;
RA Selander M., Persson E., Stenflo J., Drakenberg T.;
RT "1H NMR assignment and secondary structure of the Ca2(+)-free form of
RT the amino-terminal epidermal growth factor like domain in coagulation
RT factor X.";
RN Biochemistry 29:8111-8118(1990).
RN [11]
RX STRUCTURE BY NMR OF 85-126.
RA MEDLINE=92329412; PubMed=1627540;
RA Ullner M., Selander M., Persson E., Stenflo J., Drakenberg T.,
RA Teleman O.;
RT "Three-dimensional structure of the apo form of the N-terminal
RT EGF-like module of blood coagulation factor X as determined by NMR
RT spectroscopy and simulated folding.";
RN Biochemistry 31:5974-5983(1992).
RN [12]
RX STRUCTURE BY NMR OF 85-126.
RA MEDLINE=92406922; PubMed=1527084;
RA Selander M., Ullner M., Persson E., Teleman O.,
RA Stenflo J., Drakenberg T.;
RT "How an epidermal growth factor (EGF)-like domain binds calcium. High
RT resolution NMR structure of the calcium form of the NH2-terminal EGF-
RT like domain in coagulation factor X.";
RN J. Biol. Chem. 267:19642-19649(1992).
RN [13]
RX STRUCTURE BY NMR OF 41-126.
RA MEDLINE=96387194; PubMed=8794734;
RA Sunnerhagen M., Olah G.A., Stenflo J., Forsen S., Drakenberg T.,
RA Tremhella J.;
RT "The relative orientation of Gla and EGF domains in coagulation
RT factor X is altered by Ca2+ binding to the first EGF domain. A
RT combined NMR-small angle X-ray scattering study.";
RN Biochemistry 35:11547-11559(1996).
RN [14]
RX FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that
RX converts prothrombin to thrombin in the presence of factor Va,
RX calcium and phospholipid during blood clotting.
RX CATALYTIC ACTIVITY: Preferential cleavage: Arg-1-Thr and then
RX Arg-1-Ile bonds in prothrombin to form thrombin.
RX SUBUNIT: THE TWO CHAINS ARE FORMED FROM A SINGLE-CHAIN PRECURSOR
RX BY THE EXCISION OF TWO ARG RESIDUES AND ARE HELD TOGETHER BY 1 OR

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FT  MOD_RES 56 56 GAMMA-CARBOXYGLUTAMIC ACID.
FT  MOD_RES 59 59 GAMMA-CARBOXYGLUTAMIC ACID.
FT  MOD_RES 60 60 GAMMA-CARBOXYGLUTAMIC ACID.
FT  MOD_RES 65 65 GAMMA-CARBOXYGLUTAMIC ACID.
FT  MOD_RES 66 66 GAMMA-CARBOXYGLUTAMIC ACID.

Query Match 54.34; Score 108; DB 1; Length 492;
Best Local Similarity 40.94; Pred. No. 4; Be-11;
Matches 18; Conservative 9; Mismatches 17; Indels 0; Gaps

OY 1 ANSFLXXLRHGSLRXRCIXICXIDCFXXAKXIF2VDVDDTLAEPMSKH 44
Db 41 ANSFLVEYKQGNLERECLEFACSLFEAREVEEDAQIDDEFMSKY 84

RESULT 8
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ID PA10_HUMAN
AC P00742; Q14340; 01, Created
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 01-OCT-1989 (Rel. 12, Last annotation update)
DT 15-JUN-2002 (Rel. 11, Last annotation update)
DE Coagulation factor X precursor (EC 3.4.21.6) (Stuart factor).
GN F10.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_Taxid=9606;
RN 11
RP SEQUENCE FROM N.A.
RX MEDLINE=91216473; PubMed=1902434;
RA Messler T.L., Pittman D.D., Long G.L., Kaufman R.J., Church W.R.;
RT "Cloning and expression in COS-1 cells of a full-length cDNA encoding
RT human coagulation factor X."
RL Gene 99:291-294(1991).
RN 12
RP SEQUENCE FROM N.A.
RX MEDLINE=87026600; PubMed=3766336;
RA Leytus S.P., Foster D.C., Kurachi K., Davie E.W.;
RT "Gene for human factor X: a blood coagulation factor whose gene
RT organization is essentially identical with that of factor IX and
RT protein C."
RL Biochemistry 25:5098-5102(1986).
RN 13
RP SEQUENCE OF 13-488 FROM N.A.
RX MEDLINE=85216545; PubMed=2582420;
RA Pung M.R., Hay C.W., McGallivray R.T.A.;
RT "Characterization of an almost full-length cDNA coding for human
RT blood coagulation factor X."
RL Proc. Natl. Acad. Sci. U.S.A. 82:3591-3595(1985).
RN 14
RP SEQUENCE OF 19-488 FROM N.A.
RX TISSUE=Liver.
RX MEDLINE=86221713; PubMed=3011603;
RA Kaul R.K., Hildebrand B., Roberts S., Jagadeeswaran P.;
RT "Isolation and characterization of human blood-coagulation factor X
RT cDNA."
RL Gene 41:311-314(1986).
RN 15
RP SEQUENCE OF 41-179.
RX MEDLINE=83257207; PubMed=6871167;
RA Mcullen B.A., Fujikawa K., Kisiel W., Sasagawa T., Howard W.N.,
RA Kwa E.Y., Weinstein B.;
RT "Complete amino acid sequence of the light chain of human blood
RT coagulation factor X: evidence for identification of residue 63 as
RT beta-hydroxyaspartic acid."
RL Biochemistry 22:2875-2884(1983).
RN 16
RP SEQUENCE OF 115-488 FROM N.A., AND TISSUE SPECIFICITY.
RX TISSUE=Liver.
RX MEDLINE=84222026; PubMed=5587384;
RA Leytus S.P., Chung D.W., Kisiel W., Kurachi K., Davie E.W.;
RT "Characterization of a cDNA coding for human factor X."

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RL Proc. Natl. Acad. Sci. U.S.A. 81:3699-3702(1984).  
 RN [7]  
 RP SEQUENCE OF 183-234, AND CARBOHYDRATE-LINKAGE SITES.  
 RA MEDLINE=94062825; PubMed=8243461;  
 RX Inoue K., Morita T.;  
 RT "Identification of O-linked oligosaccharide chains in the activation  
 RT peptides of blood coagulation factor X. The role of the carbohydrate  
 RT moieties in the activation of factor X.";  
 RL Eur. J. Biochem. 218:153-163(1993).  
 RN [8]  
 RP SEQUENCE OF 1-23 FROM N.A.  
 RA MEDLINE=90128299; PubMed=2612918;  
 RX Jagadeeswaran P., Reddy S.V., Rao K.J., Hamsabhusanam K., Lyman G.;  
 RT "Cloning and characterization of the 5' end (exon 1) of the gene  
 RT encoding human factor X.";  
 RL Gene 84:517-519(1989).  
 RN [9]  
 RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS) OF 86-179 AND 235-278.  
 RX MEDLINE=93360277; PubMed=8355279;  
 RA Padmanabhan K., Padmanabhan K.P., Tulinsky A., Park C.H., Bode W.,  
 RT Huber R., Blankenship D.T., Cardin A.D., Kistiel W.;  
 RL "Structure of human des(1-45) factor Xa at 2.2-A resolution.";  
 RL J. Mol. Biol. 233:947-966(1993).  
 RN [10]  
 RP X-RAY CRYSTALLOGRAPHY (2.3 ANGSTROMS) OF 86-179 AND 235-278.  
 RX MEDLINE=98283982; PubMed=9618463;  
 RA Kamata K., Kawamoto H., Honma T., Iwama T., Kim S.H.;  
 RT "Structural basis for chemical inhibition of human blood coagulation  
 RT factor Xa.";  
 RL Proc. Natl. Acad. Sci. U.S.A. 95:6630-6635(1998).  
 CC -I- FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that  
 CC converts prothrombin to thrombin in the presence of factor Va,  
 CC calcium and phospholipid during blood clotting.  
 CC -I- CATALYTIC ACTIVITY: Preferential cleavage: Arg-|-Thr and then  
 CC Arg-|-Ile bonds in prothrombin to form thrombin.  
 CC -I- SUBUNIT: THE TWO CHAINS ARE FORMED FROM A SINGLE-CHAIN PRECURSOR  
 CC BY THE EXCISION OF TWO ARG RESIDUES AND ARE HELD TOGETHER BY 1 OR  
 CC MORE DISULFIDE BONDS.  
 CC -I- TISSUE SPECIFICITY: Plasma; synthesized in the liver.  
 CC -I- PTM: THE VITAMIN K-DEPENDENT, ENZYMATIC CARBOXYLATION OF SOME  
 CC GLUTAMIC ACID RESIDUES ALLOWS THE MODIFIED PROTEIN TO BIND  
 CC CALCIUM.  
 CC -I- PTM: N- AND O-GLYCOSYLATED.  
 CC -I- PTM: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE  
 CC INTRINSIC PATHWAY), OR BY FACTOR VIIA (IN THE EXTRINSIC PATHWAY).  
 CC -I- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.  
 CC -I- SIMILARITY: CONTAINS 2 EGF-LIKE DOMAINS.  
 CC -----  
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration  
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 CC -----  
 DR EMBL: K03194; AAA52490.1; -;  
 DR EMBL: M57285; AAA52421.1; -;  
 DR EMBL: L29433; AAA52764.1; -;  
 DR EMBL: L00390; AAA52764.1; JOINED.  
 DR EMBL: L00391; AAA52764.1; JOINED.  
 DR EMBL: L00392; AAA52764.1; JOINED.  
 DR EMBL: L00393; AAA52764.1; JOINED.  
 DR EMBL: L00394; AAA52764.1; JOINED.  
 DR EMBL: L00395; AAA52764.1; JOINED.  
 DR EMBL: L00396; AAA52764.1; JOINED.  
 DR EMBL: M2613; AAA51984.1; -;  
 DR EMBL: K01866; AAA52486.1; -;  
 DR EMBL: M33297; AAA52636.1; -;  
 DR PIR: A00924; EXHU;  
 DR PIR: A25853; A25853;  
 DR PIR: A24478; A24478;  
 DR PDB: 1HCG; 08-MAY-95.

DR PDB: 1FAX; 29-OCT-97.  
 DR PDB: 1EXY; 17-JUN-98.  
 DR PDB: 1XKA; 23-MAR-99.  
 DR PDB: 1XKB; 23-MAR-99.  
 DR MEROPS: S01\_216; -;  
 DR GlycoSuiteDB: P00742; -;  
 DR Genew: HGNC:3528; F10.  
 DR MIM: 134530; -;  
 DR MIM: 227600; -;  
 DR InterPro: IPR000152; Asx\_hydroxyl.  
 DR InterPro: IPR001314; Chymotrypsin.  
 DR InterPro: IPR000561; EGF-like.  
 DR InterPro: IPR000742; EGF-2.  
 DR InterPro: IPR001881; EGF-Ca.  
 DR InterPro: IPR002383; GLA\_blood.  
 DR InterPro: IPR001254; Ser\_protease\_Try.  
 DR InterPro: IPR000294; Vitk\_dep\_GLA.  
 DR Pfam: PF00008; EGF; 2.  
 DR Pfam: PF00089; trypsin; 1.  
 DR Pfam: PF00594; gla; 1.  
 DR PRINTS: PR00722; CHYMOTRYPSIN.  
 DR PRINTS: PR00001; GLABLOOD.  
 DR SMART: SM00179; EGF\_CA; 1.  
 DR SMART: SM00001; EGF\_Like; 1.  
 DR SMART: SM00069; GLA; 1.  
 DR SMART: SM00020; TRYP\_SPE; 1.  
 DR PROSITE: PS00010; ASX\_HYDROXYL; 1.  
 DR PROSITE: PS00022; EGF\_1; 1.  
 DR PROSITE: PS01186; EGF\_2; 2.  
 DR PROSITE: PS01187; EGF\_CA; 1.  
 DR PROSITE: PS00011; GLU\_CARBOXYLATION; 1.  
 DR PROSITE: PS00240; TRYPSIN\_DOM; 1.  
 DR PROSITE: PS00134; TRYPSIN\_HIS; 1.  
 DR PROSITE: PS00135; TRYPSIN\_SER; 1.  
 KW Glycoprotein. Hydrolyase. Serine protease. Plasma; Blood coagulation;  
 KW Gamma-carboxyglutamic acid; Hydroxylation; Calcium binding; Vitamin K;  
 KW Signal; Zymogen; EGF-like domain; Repeat; 3D-structure.  
 FT SIGNAL 1 31  
 FT PROPEP 32 40  
 FT CHAIN 41 179  
 FT CHAIN 183 488  
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 FT CHAIN 235 488  
 FT DOMAIN 86 122  
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 FT ACT\_SITE 276 276  
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 FT DISULFID 112 121  
 FT DISULFID 129 140  
 Query Match 53.8%; Score 107; DB 1; Length 488;



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Best Local Similarity 46.3%; Pred. No. 7,2e-10;
Matches 19; Conservative 5; Mismatches 17; Indels 0; Gaps 0

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Db 40 ANSFLEELRPGSLERCKEKLCSFEAREVFOSTERTKQFW 80
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RESULT 11
FA10_RABIT
ID FA10_RABIT STANDARD: PRT: 490 AA.
AC 019045:
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-JUN-2002 (Rel. 41, Last annotation update)
DE Cosignation factor X precursor (EC 3.4.21.6) (Stuart factor).
GN F10.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
RX NCBI_TaxID=9986;
RN [1]

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RX SEQUENCE FROM N.A. PubMed-9101642.
RA Pendurthi U.R., Anderson K.D., James H.L.;
RT "Characterization of a full-length cDNA for rabbit factor X";
RL Thromb. Res. 85:503-514(1997).
CC -1- FUNCTION: Factor Xa is a vitamin K-dependent glycoprotein that
CC converts prothrombin to thrombin in the presence of factor Va,
CC calcium and phospholipid during blood clotting.
CC -1- CATALYTIC ACTIVITY: Preferential cleavage: Arg-1-Thr and then
CC Arg-1-Ile bonds in prothrombin to form thrombin.
CC -1- SUBUNIT: THE TWO CHAINS ARE FORMED FROM A SINGLE-CHAIN PRECURSOR
CC BY THE EXCISION OF TWO ARG RESIDUES AND ARE HELD TOGETHER BY 1 OR
CC MORE DISULFIDE BONDS.
CC -1- PTR: THE VITAMIN K-DEPENDENT, ENZYMATIC CARBOXYLATION OF SOME
CC GLUTAMIC ACID RESIDUES ALLOWS THE MODIFIED PROTEIN TO BIND
CC CALCIUM (BY SIMILARITY).
CC -1- PTR: N- AND O-GLYCOSYLATED (BY SIMILARITY).
CC -1- PTR: THE ACTIVATION PEPTIDE IS CLEAVED BY FACTOR IXA (IN THE
CC INTRINSIC PATHWAY), OR BY FACTOR VIIA (IN THE EXTRINSIC PATHWAY)
CC (BY SIMILARITY).
CC -1- MISCELLANEOUS: CALCIUM ALSO BINDS, WITH STRONGER AFFINITY TO
CC ANOTHER SITE, BEYOND THE GLA DOMAIN.
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1.
CC -1- SIMILARITY: CONTAINS 2 EGF-LIKE DOMAINS.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
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CC -----
CC EMBL; AF003200; AAB62542.1; -.
CC HSSP; P00742; IHCG.
CC MEROPS; S01.216; -.
DR InterPro: IPR000152; Asx_hydroxyl.
DR InterPro: IPR001314; Chymotrypsin.
DR InterPro: IPR000561; EGF-like.
DR InterPro: IPR000742; EGF_2.
DR InterPro: IPR001881; EGF_CA.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Ser_protease_Try.
DR InterPro: IPR000294; VitK_dep_GLA.
DR Pfam; PF00008; EGF_2.
DR Pfam; PF00089; trypsin; 1.
DR Pfam; PF00594; gla; 1.
DR PRINTS; PR00722; CHYMOTRYPSIN.
DR PRINTS; PR00001; GLABLOOD.
DR SMART; SM00179; EGF_CA; 1.
DR SMART; SM00001; EGF_like; 1.
DR SMART; SM00069; GLA; 1.
DR SMART; SM00020; Tryp_spec; 1.
DR PROSITE; PS00010; ASX_HYDROXYL; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS00022; EGF_1; 1.
DR PROSITE; PS01186; EGF_2; 2.
DR PROSITE; PS01187; EGF_CA; 1.
DR PROSITE; PS00011; GLU_CARBOXYLATON; 1.
DR PROSITE; PS50240; TRYPSIN_DOM; 1.
DR PROSITE; PS00134; TRYPSIN_HIS; 1.
DR PROSITE; PS00133; TRYPSIN_SER; 1.
KW Glyco-carboxyl: Hydroxylase; Serine protease; Plasma; Blood coagulation;
KW Gamma-carboxyglutamic acid; Hydroxylation; Calcium-binding; Vitamin K;
KW Signal; zymogen; EGF-like domain; Repeat.
KW SIGNAL; 1 20 POTENTIAL.
FT PROPEP 21 40 BY SIMILARITY.
FT CHAIN 41 180 FACTOR X LIGHT CHAIN.
FT CHAIN 184 490 FACTOR X HEAVY CHAIN.
FT PROPEP 184 232 ACTIVATION PEPTIDE.
FT CHAIN 233 490 ACTIVATED FACTOR XA, HEAVY CHAIN.
FT DOMAIN 86 122 EGF-LIKE 1, CALCIUM-BINDING (POTENTIAL).
FT DOMAIN 125 165 EGF-LIKE 2.
FT DOMAIN 233 490 SERINE PROTEASE.

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FT MOD_RES 46 46 GAMMA-CARBOXYGLUTAMIC ACID (BY
FT MOD_RES 47 47 SIMILARITY).
FT MOD_RES 47 47 GAMMA-CARBOXYGLUTAMIC ACID (BY
FT MOD_RES 54 54 SIMILARITY).
FT MOD_RES 54 54 GAMMA-CARBOXYGLUTAMIC ACID (BY
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FT MOD_RES 79 79 SIMILARITY).
FT MOD_RES 79 79 GAMMA-CARBOXYGLUTAMIC ACID (BY
FT MOD_RES 103 103 SIMILARITY).
FT ACT_SITE 274 103 HYDROXYLATION (BY SIMILARITY).
FT ACT_SITE 320 274 CHARGE RELAY SYSTEM.
FT ACT_SITE 417 320 CHARGE RELAY SYSTEM.
FT DISULFID 90 101 CHARGE RELAY SYSTEM.
FT DISULFID 95 110 BY SIMILARITY.
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FT DISULFID 129 140 BY SIMILARITY.
FT DISULFID 136 149 BY SIMILARITY.
FT DISULFID 151 164 BY SIMILARITY.
FT DISULFID 172 340 INTERCHAIN (BY SIMILARITY).
FT DISULFID 239 244 BY SIMILARITY.
FT DISULFID 259 275 BY SIMILARITY.
FT DISULFID 388 402 BY SIMILARITY.
FT DISULFID 413 441 BY SIMILARITY.
FT CARBOHYD 61 61 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 187 187 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 205 205 N-LINKED (GLCNAC. . .) (POTENTIAL).
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Query Match 50.3%: Score 100: DB 1: Length 490:
Best Local Similarity 40.9%: Pred: 1.2e-09:
Matches 18: Conservative 9: Mismatches 17: Indels 0: Gaps 0:

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RA Davie E.W.;
RA "Characterization of a cDNA coding for human factor VII.";
RA Proc. Natl. Acad. Sci. U.S.A. 83:2412-2416(1986).
RL [2]
RP SEQUENCE FROM N.A.
RX MEDLINE-87260948: PubMed-3037537;
RA O'Hara P.J., Grant F.J., Haldeman B.A., Gray C.L., Insley M.Y.,
RA Hagen F.S., Murray M.J.;
RA "Nucleotide sequence of the gene coding for human factor VII, a
RA vitamin K-dependent protein participating in blood coagulation.";
RA Proc. Natl. Acad. Sci. U.S.A. 84:5158-5162(1987).
RL [3]
RN SEQUENCE FROM N.A., AND VARIANTS THR-352; GLN-413 AND LYS-445.
RA Rieder M.J., Armet T.Z., Carrington D.P., Chung M.-W., Lee K.L.,
RA Poel C.L., Toth E.J., Yi Q., Nickerson D.A.;
RA Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
RN [4]
RX SEQUENCE OF 61-466, AND POST-TRANSLATIONAL MODIFICATIONS.
RX MEDLINE-89088153: PubMed-3264725;
RA Thim L., Bjoern S., Christensen M., Nicolaisen E.M., Lund-Hansen T.,
RA Pedersen A.H., Hedner U.;
RA "Amino acid sequence and posttranslational modifications of human
RA factor VIIa from plasma and transfected baby hamster kidney cells.";
RA Biochemistry 27:7785-7793(1988).
RL [5]
RN CARBOHYDRATE-LINKAGE SITES SER-112 AND SER-120.
RX MEDLINE-91250411: PubMed-1904059;
RA Bjoern S., Foster D.C., Thim L., Wiberg F.C., Christensen M.,
RA Komiyama Y., Pedersen A.H., Kistiel W.;
RA "Human plasma and recombinant factor VII. Characterization of O-
RA glycosylations at serine residues 52 and 60 and effects of site-
RA directed mutagenesis of serine 52 to alanine.";
RA J. Biol. Chem. 266:11051-11057(1991).
RL [6]
RN STRUCTURE OF CARBOHYDRATE ON SER-112.
RX MEDLINE-90062160: PubMed-2511201;
RA Nishimura H., Kawabata S., Kistiel W., Hase S., Ikenaka T., Takao T.,
RA Shimomishi Y., Iwanaga S.;
RA "Identification of a disaccharide (Xyl-Glc) and a trisaccharide
RA (Xyl2-Glc) O-glycosidically linked to a serine residue in the first
RA epidermal growth factor-like domain of human factors VII and IX and
RA protein Z and bovine protein Z.";
RA J. Biol. Chem. 264:20320-20325(1989).
RL [7]
RN STRUCTURE OF CARBOHYDRATE ON SER-112.
RX MEDLINE-91344709: PubMed-2129367;
RA Iwanaga S., Nishimura H., Kawabata S., Kistiel W., Hase S., Ikenaka T.;
RA "A new trisaccharide sugar chain linked to a serine residue in the
RA first EGF-like domain of clotting factors VII and IX and protein Z.";
RA Adv. Exp. Med. Biol. 281:121-131(1990).
RL [8]
RN X-RAY CRYSTALLOGRAPHY (2.0 ANGSTROMS) OF FVIIA IN COMPLEX WITH TF.
RX MEDLINE-96175641: PubMed-8598903;
RA Banner D.W., D'Arcy A., Chene C., Winkler F.K., Guha A.,
RA Konigsberg W.H., Nemerson Y., Kirchhofer D.;
RA "The crystal structure of the complex of blood coagulation factor
RA VIIa with soluble tissue factor.";
RA Nature 380:41-46(1996).
RL [9]
RN X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF FVIIA IN COMPLEX WITH TF.
RX MEDLINE-99126538: PubMed-9925787;
RA Zhang E., St. Charles R., Tulinsky A.;
RA "Structure of extracellular tissue factor complexed with factor VIIa
RA inhibited with a BPTI mutant.";
RA J. Mol. Biol. 285:2089-2104(1999).
RL [10]
RN STRUCTURE BY NMR OF 105-145.
RX MEDLINE-98367502: PubMed-9692950;
RA Murayai A., Finn B.E., Gippert G.P., Forsen S., Stenflo J.,
RA Drakenberg T.;
RA "Solution structure of the N-terminal EGF-like domain from human
RA factor VII.";
RA Biochemistry 37:10605-10615(1998).

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CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).

DR EMBL: AF009242; AAB67070.1; -  
DR HSSP: P00740; 1CFH.  
DR Genew: HGNC:9469; PRRG1.  
DR MIM: 604428; -  
DR InterPro: IPR002383; GLA\_blood.  
DR InterPro: IPR000294; VitK\_dep\_GLA.  
DR Pfam: PF00594; gla; 1.  
DR PRINTS: PR00001; GLABLOOD.  
DR SMART: SM00069; GLA; 1.  
DR PROSITE: PS00011; GLU\_CARBOXYLATION; 1.  
DR Gamma-carboxyglutamic acid; Vitamin K; Transmembrane.  
KW PROPEP  
FT CHAIN 1 20  
FT TRANSMEMBRANE GAMMA-CARBOXYGLUTAMIC ACID  
FT PROTEIN 1.  
FT DOMAIN 21 83 EXTRACELLULAR (POTENTIAL).  
FT TRANSMEM 84 106 POTENTIAL.  
FT DOMAIN 107 218 CYTOPLASMIC (POTENTIAL).  
FT DOMAIN 24 61 GLA-RICH.  
FT DOMAIN 131 135 POLY-PRO.  
SQ SEQUENCE 218 AA; 24947 MW; 26538A61AB0AE98 CRC64;

Query Match 43.2%; Score 86; DB 1; Length 218;  
Best Local Similarity 34.1%; Pred. No. 1.4e-07;  
Matches 15; Conservative 8; Mismatches 21; Indels 0; Gaps 0;

OY 1 ANSFLXXLRHGSIXRCIXXCIFXAXKXIFZDVDDTLAFWSKH 44  
DB 21 ANCFEEIRGNIERCKEECTFEARAFENNEXTKFEWSTY 64

RESULT 14  
TMG4\_HUMAN STANDARD; PRT; 226 AA.  
AC 09BZD6;  
DT 15-JUN-2002 (Rel. 41, Created)  
DT 15-JUN-2002 (Rel. 41, Last sequence update)  
DT 15-JUN-2002 (Rel. 41, Last annotation update)  
DE Transmembrane gamma-carboxyglutamic acid protein 4 precursor.  
GN TMG4.  
OS Homo sapiens (Human).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
OX NCBI\_TaxID=9606;  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE-21117044; PubMed-11171957;  
RA Kulan J.D., Harris J.E., Xie L., Dayle E.W.;  
RT "Identification of two novel transmembrane gamma-carboxyglutamic acid  
RT proteins expressed broadly in fetal and adult tissues.";  
RT Proc. Natl. Acad. Sci. U.S.A. 98:1370-1375(2001).  
RN [2]  
RP SEQUENCE FROM N.A.  
RC TISSUE=Placenta;  
RL Submitted (JUL-2001) to the EMBL/Genbank/DDBJ databases.  
CC -1- SURCELLULAR LOCATION: Type I membrane protein.  
CC -1- TISSUE SPECIFICITY: Expressed in lung, liver, kidney, pancreas and  
CC placenta.  
CC -1- PTM: Gla residues are produced after subsequent posttranslational  
CC modifications of glutamic acid by a vitamin K-dependent gamma-  
CC carboxylase.  
CC This SWISS-PROT entry is copyright. It is produced through a collaboration  
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DR EMBL: AF326351; AAK0956.1; -  
DR EMBL: BC010052; AAH10052.1; -  
DR HSSP: P00740; 1CFH.  
DR InterPro: IPR002383; GLA\_blood.  
DR InterPro: IPR000294; VitK\_dep\_GLA.  
DR Pfam: PF00594; gla; 1.  
DR PRINTS: PR00001; GLABLOOD.  
DR SMART: SM00069; GLA; 1.  
DR PROSITE: PS00011; GLU\_CARBOXYLATION; 1.  
DR Gamma-carboxyglutamic acid; Vitamin K; Transmembrane; Signal.  
FT SIGNAL 1 17  
FT PROPEP 18 49  
FT CHAIN 50 226  
FT TRANSMEMBRANE GAMMA-CARBOXYGLUTAMIC ACID  
FT PROTEIN 4.  
FT DOMAIN 50 113 EXTRACELLULAR (POTENTIAL).  
FT TRANSMEM 114 134 POTENTIAL.  
FT DOMAIN 135 226 CYTOPLASMIC (POTENTIAL).  
FT DOMAIN 56 93 GLA-RICH.  
FT DOMAIN 203 208 POLY-PRO.  
SQ SEQUENCE 226 AA; 25403 MW; 45C783E3825797EE CRC64;

Query Match 42.2%; Score 84; DB 1; Length 226;  
Best Local Similarity 41.2%; Pred. No. 3.2e-07;  
Matches 14; Conservative 8; Mismatches 12; Indels 0; Gaps 0;

OY 11 GSLRXRCIXXCIFXAXKXIFZDVDDTLAFWSKH 44  
DB 63 GNLERCNEELCNVEERLEIFVEDKTIAPWQEX 96

RESULT 15  
FA7\_BOVIN STANDARD; PRT; 407 AA.  
AC P22457;  
DT 01-AUG-1991 (Rel. 19, Created)  
DT 01-AUG-1991 (Rel. 19, Last sequence update)  
DT 15-JUN-2002 (Rel. 41, Last annotation update)  
DE Coagulation factor VII (EC 3.4.21.21) (Serum prothrombin conversion  
DE accelerator).  
GN F7.  
OS Bos taurus (Bovine).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
OC Bovidae; Bovinae; Bos.  
OX NCBI\_TaxID=9913;  
RN [1]  
RP SEQUENCE.  
RX MEDLINE-89008362; PubMed-3049594;  
RA Takeya H., Kawabata S., Nakagawa K., Yamamichi Y., Miyata T.,  
RA Iwanaga S.;  
RT "Bovine factor VII. Its purification and complete amino acid  
RT sequence.";  
RT J. Biol. Chem. 263:14866-14877(1988).  
RN [2]  
RP STRUCTURE OF CARBOHYDRATE ON SER-52.  
RX MEDLINE-89213999; PubMed-3149637;  
RA Hase S., Kawabata S., Nishimura H., Takeya H., Sueyoshi T.,  
RA Miyata T., Iwanaga S., Takao T., Shimomishi Y., Ikenaka T.;  
RT "A new trisaccharide sugar chain linked to a serine residue in bovine  
RT blood coagulation factors VII and IX.";  
RT J. Biochem. 104:867-868(1988).  
RN [3]  
RP STRUCTURE OF CARBOHYDRATE ON SER-52.  
RX MEDLINE-91344709; PubMed-2129367;  
RA Iwanaga S., Nishimura H., Kawabata S., Kistel W., Hase S., Ikenaka T.;  
RT "A new trisaccharide sugar chain linked to a serine residue in the  
RT first EGF-like domain of clotting factors VII and IX and protein Z.";  
RT Adv. Exp. Med. Biol. 281:121-131(1990).



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## OM protein - protein search, using sw model

Run on: May 13, 2003, 14:01:48 : Search time 29 Seconds

(without alignments)  
312.623 Million cell updates/sec

Title: SEQ1-EDITED

Perfect score: 199

Sequence: 1 ANSFLXLRHSLKRCXIX.....XXAKXIFZVDVDTLAFMSKH 44

Scoring table: BIOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 671580 seqs, 206047115 residues

Total number of hits satisfying chosen parameters: 671580

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :  
1: SP archaea: \*  
2: SP bacteria: \*  
3: SP fungi: \*  
4: SP human: \*  
5: SP\_invertebrate: \*  
6: SP\_mammal: \*  
7: SP\_mhc: \*  
8: SP\_organelle: \*  
9: SP\_phage: \*  
10: SP\_plant: \*  
11: SP\_rodent: \*  
12: SP\_virus: \*  
13: SP\_vertebrate: \*  
14: SP\_unclassified: \*  
15: SP\_virus: \*  
16: SP\_bacteriap: \*  
17: SP\_archaeap: \*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	151	75.9	456	09TTR0	09TTR0 canis fam1
2	140	70.4	460	091WN8	091WN8 mus musculu
3	134	67.3	460	099PC6	099PC6 mus musculu
4	112	56.3	482	063207	063207 rattus norv
5	98	49.2	481	054740	054740 mus musculu
6	98	49.2	481	099L32	099L32 mus musculu
7	98	49.2	481	088947	088947 mus musculu
8	98	49.2	701	096P08	096P08 homo sapien
9	92	46.2	469	096MD9	096MD9 orlithorhyn
10	92	46.2	100	015253	015253 homo sapien
11	79	39.7	446	061109	061109 mus musculu
12	79	39.7	650	09NSD0	09NSD0 homo sapien
13	79	39.7	650	016519	016519 homo sapien
14	76.5	38.4	542	08T613	08T613 halocynthia
15	76	38.2	138	028994	028994 sus scrofa
16	75	37.7	607	091001	091001 gallus gall

17	74	37.2	456	4	014316	014316 homo sapien
18	74	37.2	461	6	095ND7	095ND7 pan troglod
19	74	37.2	461	6	095ND6	095ND6 pan troglod
20	72	36.2	648	6	029094	029094 sus scrofa
21	71.5	35.9	433	13	0290YK1	0290YK1 brachydanto
22	71	35.7	49	6	095ME8	095ME8 bos taurus
23	70	35.2	98	13	P82807	P82807 notechlis sc
24	69	34.7	608	13	09PTW7	09PTW7 struthio ca
25	67	33.7	399	11	09COM3	09COM3 mus musculu
26	64	32.2	25	11	09OVH6	09OVH6 rattus sp.
27	63	31.7	179	4	08TAS3	08TAS3 homo sapien
28	63	31.7	198	11	08R182	08R182 mus musculu
29	59	29.6	673	11	061592	061592 mus musculu
30	58	29.6	674	11	099K57	099K57 mus musculu
31	58	29.1	674	11	063772	063772 rattus sp.
32	57	28.6	678	4	Q14393	Q14393 homo sapien
33	56.5	28.4	606	10	09SUG9	09SUG9 arabidopsis
34	56.5	28.4	651	10	08S218	08S218 oryza sativ
35	55.5	27.9	459	10	09SE22	09SE22 oryza sativ
36	55.5	27.9	575	10	094E17	094E17 oryza sativ
37	54.5	27.4	603	10	09LPG7	09LPG7 arabidopsis
38	53.5	26.9	196	10	004284	004284 selaginella
39	53.5	26.9	567	10	08W4J2	08W4J2 arabidopsis
40	53.5	26.9	593	10	09LUC3	09LUC3 arabidopsis
41	53.5	26.9	608	10	09XF36	09XF36 medicago sa
42	52.5	26.4	431	10	094EY5	094EY5 arabidopsis
43	52.5	26.4	506	10	09SPF0	09SPF0 oryza sativ
44	52.5	26.4	506	10	09SE23	09SE23 oryza sativ
45	52.5	26.4	543	10	Q9MB23	Q9MB23 arabidopsis

## ALIGNMENTS

## RESULT 1

ID	Q9TTR0	PRELIMINARY:	PRT:	456 AA.
AC	Q9TTR0			
DT	01-MAY-2000 (TREMBLrel. 13, Created)			
DT	01-MAY-2000 (TREMBLrel. 13, Last sequence update)			
DT	01-MAR-2002 (TREMBLrel. 20, Last annotation update)			
DE	Protein C precursor.			
GN	PROC.			
OS	Canis familiaris (Dog).			
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.			
OX	NCBI_TaxID=9615;			
RN	(1)			
RP	SEQUENCE FROM N.A.			
RA	Leeb T., Kopp T., Deppe A., Breen M., Matls U., Brunberg L.,			
RA	Brenig B.;			
RT	"Molecular characterization and chromosomal assignment of the canine			
RT	protein C gene."			
RL	Mamm. Genome 10:135-139(1999).			
RN	(2)			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=99371952; PubMed=10443005;			
RA	Leeb T., Pfeiffer I., Kopp T., Deppe A., Brenig B.;			
RA	Anim. Genet. 30:237-238(1999).			
CC	-1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE			
CC	TRYPSIN FAMILY.			
CC	EMBL: AJ001979; CAA05126.1; -			
DR	HSSP: P04070; IPCU.			
DR	MEROPS: S01.218; -			
DR	InterPro: IPR000152; Asx_hydroxyl.			
DR	InterPro: IPR001314; Chymotrypsin.			
DR	InterPro: IPR000561; EGF-like.			
DR	InterPro: IPR001881; EGF Ca.			
DR	InterPro: IPR002383; GLA_blood.			
DR	InterPro: IPR001254; Ser_protease_Try.			
DR	InterPro: IPR000294; Vitr_dep_GLA.			
DR	Pfam: PF00008; EGF; 2.			

DR Pfam: PF00594; gla; 1.  
 DR Pfam: PF00089; trypsin; 1.  
 DR PRINTS: PR00722; CHYMOTRYPSIN.  
 DR PRINTS: PR00001; GLABLOOD.  
 DR SMART: SM00181; EGF; 2.  
 DR SMART: SM00069; GLA; 1.  
 DR SMART: SM00020; TRYP\_SPE; 1.  
 DR PROSITE: PS00010; ASX\_HYDROXYL; 1.  
 DR PROSITE: PS00022; EGF\_1; UNKNOWN\_1.  
 DR PROSITE: PS01186; EGF\_2; 2.  
 DR PROSITE: PS01187; EGF\_CA; 1.  
 DR PROSITE: PS00011; GLU\_CARBOXYLATION; 1.  
 DR PROSITE: PS50240; TRYPsin\_DOM; 1.  
 DR PROSITE: PS00134; TRYPsin\_HIS; UNKNOWN\_1.  
 DR PROSITE: PS00135; TRYPsin\_SER; 1.  
 DR Calcium-binding: EGF-like domain; Glycoprotein; Hydrolase;  
 KW Hydroxylation; Repeat; Serine protease; Signal.  
 FT SIGNAL 1 42  
 FT CHAIN 43 192 PROTEIN C LIGHT CHAIN.  
 FT CHAIN 193 194 PROTEIN C CONNECTING DIPEPTIDE.  
 FT CHAIN 195 456 PROTEIN C HEAVY CHAIN.  
 SQ SEQUENCE 456 AA; 50813 MW; 7AD3A8C1C34E59FF CRC64;

Query Match 75.9%; Score 151; DB 6; Length 456;  
 Best Local Similarity 63.6%; Pred. No. 1.2e-17;  
 Matches 28; Conservative 6; Mismatches 10; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSIXRCIXICDFXAXKXIFZDVDDTLAFWSKH 44  
 DB 43 ANSFLFETIRAGSLERECMEICDFEPAKEIFONVEDTLAFWISKY 86

## RESULT 2

ID 091WN8 PRELIMINARY; PRT; 460 AA.  
 AC 091WN8;  
 DT 01-DEC-2001 (TRENBLREL. 19, Created)  
 DT 01-DEC-2001 (TRENBLREL. 19, Last sequence update)  
 DT 01-JUN-2002 (TRENBLREL. 21, Last annotation update)  
 DE Similar to protein C.  
 GN PROC.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 OX NCBI\_Taxid=10090;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=LIVER;  
 RA Strausberg R.;  
 RL Submitted (SEP-2001) to the EMBL/Genbank/DBJ databases.  
 DR EMBL; BC013896; AAH13896.1; -  
 DR MGI; MGI:97771; Proc.  
 DR InterPro: IPR000152; Asx\_hydroxyl.  
 DR InterPro: IPR000561; EGF-like.  
 DR InterPro: IPR001881; EGF\_CA.  
 DR InterPro: IPR001254; Ser\_protease\_Try.  
 DR InterPro: IPR000294; VltK\_dep\_GLA.  
 DR Pfam: PF00008; EGF; 2.  
 DR Pfam: PF00594; gla; 1.  
 DR Pfam: PF00089; trypsin; 1.  
 DR PROSITE: PS00010; ASX\_HYDROXYL; UNKNOWN\_1.  
 DR PROSITE: PS00022; EGF\_1; UNKNOWN\_1.  
 DR PROSITE: PS01186; EGF\_2; UNKNOWN\_2.  
 DR PROSITE: PS01187; EGF\_CA; UNKNOWN\_1.  
 DR PROSITE: PS00011; GLU\_CARBOXYLATION; UNKNOWN\_1.  
 DR PROSITE: PS50240; TRYPsin\_DOM; 1.  
 DR PROSITE: PS00134; TRYPsin\_HIS; UNKNOWN\_1.  
 DR PROSITE: PS00135; TRYPsin\_SER; UNKNOWN\_1.  
 KW Hydrolase; Serine protease.  
 SQ SEQUENCE 460 AA; 51818 MW; 0117F26E68FCC274 CRC64;

Query Match 70.4%; Score 140; DB 11; Length 460;  
 Best Local Similarity 59.1%; Pred. No. 9.6e-16;

Matches 26; Conservative 7; Mismatches 11; Indels 0; Gaps 0;  
 OY 1 ANSFLXLRHGSIXRCIXICDFXAXKXIFZDVDDTLAFWSKH 44  
 DB 42 ANSFLFETIRAGSLERECMEICDFEPAKEIFONVEDTLAFWISKY 85

## RESULT 3

ID 099PC6 PRELIMINARY; PRT; 460 AA.  
 AC 099PC6;  
 DT 01-JUN-2001 (TRENBLREL. 17, Created)  
 DT 01-JUN-2001 (TRENBLREL. 17, Last sequence update)  
 DT 01-JUN-2002 (TRENBLREL. 21, Last annotation update)  
 DE Anticoagulant protein C.  
 GN PROC.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 OX NCBI\_Taxid=10090;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=C57BL;  
 RA Korf I.;  
 RT "Complete sequence of UC72A01.";  
 RL Submitted (NOV-2000) to the EMBL/Genbank/DBJ databases.  
 CC -1 SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE  
 CC TRYPSIN FAMILY.  
 CC EMBL; AF318182; AKK07918.1; -  
 DR HSSP; P04070; 1PCU.  
 DR MEROPS; S01.218; -  
 DR MGI; MGI:97771; Proc.  
 DR InterPro: IPR000152; Asx\_hydroxyl.  
 DR InterPro: IPR001314; Chymotrypsin.  
 DR InterPro: IPR000561; EGF-like.  
 DR InterPro: IPR001881; EGF\_CA.  
 DR InterPro: IPR002383; GLA\_blood.  
 DR InterPro: IPR001254; Ser\_protease\_Try.  
 DR InterPro: IPR000294; VltK\_dep\_GLA.  
 DR Pfam: PF00008; EGF; 2.  
 DR Pfam: PF00594; gla; 1.  
 DR Pfam: PF00089; trypsin; 1.  
 DR PRINTS: PR00722; CHYMOTRYPSIN.  
 DR PRINTS: PR00001; GLABLOOD.  
 DR SMART: SM00181; EGF; 2.  
 DR SMART: SM00001; EGF\_like; 2.  
 DR SMART: SM00069; GLA; 1.  
 DR SMART: SM00020; TRYP\_SPE; 1.  
 DR PROSITE: PS00010; ASX\_HYDROXYL; 1.  
 DR PROSITE: PS00022; EGF\_1; UNKNOWN\_1.  
 DR PROSITE: PS01186; EGF\_2; 2.  
 DR PROSITE: PS01187; EGF\_CA; 1.  
 DR PROSITE: PS00011; GLU\_CARBOXYLATION; 1.  
 DR PROSITE: PS50240; TRYPsin\_DOM; 1.  
 DR PROSITE: PS00134; TRYPsin\_HIS; UNKNOWN\_1.  
 DR PROSITE: PS00135; TRYPsin\_SER; 1.  
 KW Calcium-binding: EGF-like domain; Glycoprotein; Hydrolase;  
 KW Hydroxylation; Repeat; Serine protease.  
 SQ SEQUENCE 460 AA; 51784 MW; 0293BC25E9D3ED16 CRC64;

Query Match 67.3%; Score 134; DB 11; Length 460;  
 Best Local Similarity 56.8%; Pred. No. 1.1e-14;  
 Matches 25; Conservative 7; Mismatches 12; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSIXRCIXICDFXAXKXIFZDVDDTLAFWSKH 44  
 DB 42 ANSFLFETIRAGSLERECMEICDFEPAKEIFONVEDTLAFWISKY 85

## RESULT 4

ID 063207 PRELIMINARY; PRT; 482 AA.  
 AC 063207;

DT	01-NOV-1996 (Tremblrel. 01, Created)
DT	01-NOV-1996 (Tremblrel. 01, Last sequence update)
DT	01-MAR-2002 (Tremblrel. 20, Last annotation update)
DE	Factor X.
DE	Rattus norvegicus (Rat).
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX	NCBI_TaxID=10116;
RN	[1]
RP	SEQUENCE FROM N.A.
RC	STRAIN=SPRAGUE-DAWLEY;
RX	MEDLINE=96093366; PubMed=8578539;
RA	Stanton C., Ross R.P., Hutson S., Wallin R.;
RT	"Evidence for competition between vitamin K-dependent clotting factors
RT	for intracellular processing by the vitamin K-dependent gamma-
RL	carboxylase."
RL	Thromb. Res. 80:63-73(1995).
CC	-1 SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
CC	TRYPSIN FAMILY.
CC	EMBL: X79807; CAA56202.1; -.
DR	HSSP; P00742; 1XKA.
DR	MEROPS; S01.216; -.
DR	InterPro: IPR000152; Asx_hydroxyl.
DR	InterPro: IPR001314; Chymotrypsin.
DR	InterPro: IPR000561; EGF-like.
DR	InterPro: IPR000742; EGF_2.
DR	InterPro: IPR001881; EGF_Ca.
DR	InterPro: IPR002383; GLA_blood.
DR	InterPro: IPR001254; Ser_protase-Try.
DR	InterPro: IPR000294; VitK_dep_GLA.
DR	Pfam; PF00008; EGF_2.
DR	Pfam; PF00594; gla; 1.
DR	Pfam; PF00089; trypsin; 1.
DR	PRINTS; PR00722; CHYMOTRYPSIN.
DR	PRINTS; PR00001; GLABLOOD.
DR	SMART; SM00179; EGF_Ca; 1.
DR	SMART; SM00001; EGF_Like; 1.
DR	SMART; SM00059; GLA; 1.
DR	SMART; SM00020; Tryp_Spc; 1.
DR	PROSITE; PS00010; ASX_HYDROXYL; 1.
DR	PROSITE; PS00022; EGF_1; UNKNOWN_1.
DR	PROSITE; PS01186; EGF_2; 2.
DR	PROSITE; PS01187; EGF_CA; 1.
DR	PROSITE; PS00011; GLU_CARBOXYLATION; 1.
DR	PROSITE; PS02040; TRYPSIN_DOM; 1.
DR	PROSITE; PS00134; TRYPSIN_HIS; UNKNOWN_1.
DR	PROSITE; PS00135; TRYPSIN_SER; 1.
KW	Calcium-binding; EGF-like domain; Glycoprotein; Hydrolase;
KW	Hydroxylation; Repeat; Serine protease.
SO	SEQUENCE 482 AA; 54265 MW; 0284678E3954A698 CRC64;
Query Match	56.3%; Score 112; DB 11; Length 482;
Best Local Similarity	40.9%; Pred. No. 7, 5e-11;
Matches 18; Conservative 10; Mismatches 16; Indels 0; Gaps 0;	
OY	1 ANSFLXLRHGSGLARXCIXIXICDPXAKXIFZVDVDTLAFWSKH 44
DB	41 ANSFPEETIKGNLRECEVEICSPFEAREVFEDNKETIEFMNKY 84
RESULT 5	
ID	054740 PRELIMINARY; PRT; 481 AA.
AC	054740;
DT	01-JUN-1998 (Tremblrel. 06, Created)
DT	01-JUN-1998 (Tremblrel. 06, Last sequence update)
DT	01-JUN-2002 (Tremblrel. 21, Last annotation update)
DE	Coagulation factor X precursor (EC 3.4.21.6).
GN	F10 OR FA10.
OS	Mus musculus (Mouse).
OC	Plasmid plasmascapit.
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

Ox	[1]	NCBI_TaxID=10090;
Rn		
Rp		SEQUENCE FROM N.A.
Rc		TISSUE=LIVER:
Rx		MEDLINE=98454993; PubMed=9783672;
Ra		"Heidmann H.H., Kontermann R.E.;"
Rt		"Cloning and recombinant expression of mouse coagulation factor X.";
Rl		Thromb. Res. 92:33-41(1998).
Cc	-i-	SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
Cc	-	TRYPSIN FAMILY.
Dd	EMBL:	AJ222677; CAA10933.1; -
Dd	HSPB:	P00742; IYKA.
Dd	MERO:	S01.216; -
Dd	MED:	MGI:103107; F10.
Dd	InterPro:	IPR000152; Asx_hydroxyl.
Dd	InterPro:	IPR001314; Chymotrypsin.
Dd	InterPro:	IPR000561; EGF-like.
Dd	InterPro:	IPR000742; EGF_2.
Dd	InterPro:	IPR001881; EGF_Ca.
Dd	InterPro:	IPR002383; GLA_blood.
Dd	InterPro:	IPR001254; Ser_protease TRY.
Dd	InterPro:	IPR000294; Vltk_dep_GLA.
Dd	Pfam:	PF00594; gla; 1.
Dd	Pfam:	PF00089; trypsin; 1.
Dd	PRINTS:	PRO0722; CHYMOTRYPSIN.
Dd	PRINTS:	PRO0001; GLABLOOD.
Dd	SMART:	SMO0179; EGF_CA; 1.
Dd	SMART:	SMO0001; EGF_like; 1.
Dd	SMART:	SMO0069; GLA; 1.
Dd	SMART:	SMO0030; Tryp_Spc; 1.
Dd	PROSITE:	PS00010; ASX_HYDROXYL; UNKNOWN_1.
Dd	PROSITE:	PS00022; EGF_1; UNKNOWN_1.
Dd	PROSITE:	PS01186; EGF_2; 2.
Dd	PROSITE:	PS01187; EGF_CA; 1.
Dd	PROSITE:	PS00011; GLU CARBOXYLATION; 1.
Dd	PROSITE:	PS50240; TRYP SIN_DOM; 1.
Dd	PROSITE:	PS00134; TRYP SIN_HIS; UNKNOWN_1.
Dd	PROSITE:	PS00135; TRYP SIN_SER; 1.
Kw	Calcium-binding:	EGF-like domain; Glycoprotein; Hydrolase; Plasmid;
Kw	Repeat:	Serine protease; Signal.
Ft	SIGNAL	1 40 POTENTIAL.
Ft	CHAIN	41 481 COAGULATION FACTOR X
So	SEQUENCE	481 AA; 53986 MW; CF702DE5EF9D97AE CRC64;
	Query Match	49.2%; Score 98; DB 11; Length 481;
	Best Local Similarity	36.4%; Pred. No. 2e-08;
	Matches 16;	Conservative 10; Mismatches 18; Indels 0; Gaps 0.
Oy	1	ANSFLXLRHGS LKRXCIIXICDFYXAXXIFZVDVDTLAFMSKH 44       :      :     :     :
Db	41	ANSPFEFFKKCNLERECMEHCISCYEVRIRIFEDDEKTKREYWIKY 84
RESULT 6		
ID	O99L32	PRELIMINARY; PRT; 481 AA.
AC	O99L32:	
Dt	01-JUN-2001	(TREMBLrel. 17, Created)
Dt	01-JUN-2001	(TREMBLrel. 17, Last sequence update)
Dt	01-JUN-2002	(TREMBLrel. 21, last annotation update)
De	Coagulation factor X.	
Gn	F10.	
Os	Mus musculus (Mouse);	
Oc	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
Oc	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.	
Ox	NCBI_TaxID=10090;	
Rn	[1]	
Rp	SEQUENCE FROM N.A.	
Rc	Straussberg R.;	
Rl	Submitted (Feb-2001) to the EMBL/Genbank/DBJ databases.	
Cc	-i-	SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
Cc	-	TRYPSIN FAMILY.

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DR EMBL: BC003877; AA03877.1; -
DR HSSP: P00742; 1XKA.
DR MEROPS: S01_216; -
DR MGD; MG1_103107; F10.
DR InterPro: IPR000152; Asx_hydroxyl.
DR InterPro: IPR001314; Chymotrypsin.
DR InterPro: IPR000561; EGF-like.
DR InterPro: IPR000742; EGF_2.
DR InterPro: IPR001881; EGF_Ca.
DR InterPro: IPR001438; EGF_II.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Ser-protease_Try.
DR InterPro: IPR000294; Vltk_dep_GLA.
DR Pfam: PF00008; EGF_2.
DR Pfam: PF00594; gla_1.
DR Pfam: PF00089; trypsin_1.
DR PRINTS: PR00722; CHYMOTRYPSIN.
DR PRINTS: PR00010; EGFBL00D.
DR PRINTS: PR00001; GLABLOOD.
DR SMART: SM00181; EGF_2.
DR SMART: SM00001; EGF-like; 2.
DR SMART: SM00069; GLA_1.
DR SMART: SM00020; TRYP_SPE; 1.
DR PROSITE: PS00010; ASX_HYDROXYL; UNKNOWN_1.
DR PROSITE: PS00022; EGF_1; UNKNOWN_1.
DR PROSITE: PS01186; EGF_2; 2.
DR PROSITE: PS01187; EGF_CA; 1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE: PS50240; TRYPsin_DOM; 1.
DR PROSITE: PS00134; TRYPsin_HIS; UNKNOWN_1.
DR PROSITE: PS00135; TRYPsin_SER; 1.
DR Calcium-binding; EGF-like domain; Glycoprotein; Hydrolase; Repeat;
KW Serine protease.
SQ SEQUENCE 481 AA; 54004 MW; BD8BE96C8A0B7E7F CRC64;

Query Match 49.2%; Score 98; DB 11; Length 481;
Best Local Similarity 36.4%; Pred. NO. 2e-08;
Matches 16; Conservative 10; Mismatches 18; Indels 0; Gaps 0.

QY 1 ANSFLXLRHGSGLKRCXIXICDFFXAKXIFZVDVDTLAWSKH 44
    1111 1:1 1: 11: 1:1:1: 1:1:1:
DB 41 ANSFEEFRKGNLERECMEICSYEEVREIFEDDEKTKRYWKY 84

RESULT 7
ID 088947 PRELIMINARY: PRT; 481 AA.
AC 088947;
DT 01-NOV-1998 (TREMBLrel. 08, Created)
DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)
DT 01-MAR-2002 (TREMBLrel. 20, Last annotation update)
DE Coagulation factor X precursor.
GN F10.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
PI [1]
PI SEQUENCE FROM N.A.
RC STRAIN=C57BL/6 X CBA; TISSUE=LIVER;
RX MEDLINE=98347933; PubMed=9684791;
RA Liang Z., Cooper A., Deford M.E., Carmeliet P., Collen D.,
RA Castellino F.J., Rosen E.D.;
RT "Cloning and characterization of a cDNA encoding murine coagulation
RT factor X."
RL Thromb. Haemost. 80:87-91(1998).
RN [2]
RN SEQUENCE FROM N.A.
RC STRAIN=129S/J;
RA Cooper A., Liang Z., Castellino F.J., Rosen E.D.;
RT "Cloning and Characterization of the Murine Factor X Gene."
RT Thromb. Haemost. 0:0-0(2000).
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE

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CC	TRYPSIN FAMILY.
DR EMBL: AF087644; AAC36345.1; "	
DR EMBL: AF211347; AAF22980.1; "	
DR HSSP: P00742; 1KKA.	
DR MEROPS: S01.216; "	
DR MGD: MGI:103107; F10.	
DR InterPro: IPR000152; Asx_hydroxyl.	
DR InterPro: IPR001314; Chymotrypsin.	
DR InterPro: IPR000561; EGF-like.	
DR InterPro: IPR000742; EGF_2.	
DR InterPro: IPR001881; EGF_Ca.	
DR InterPro: IPR002383; GLA blood.	
DR InterPro: IPR001254; Ser_protease_Try.	
DR InterPro: IPR000294; Vltk_dep_GLA.	
DR Pfam: PF00008; EGF_2.	
DR Pfam: PF00594; gla; 1.	
DR Pfam: PF00589; trypsin; 1.	
DR PRINTS: PR00722; CHYMOTRYPSIN.	
DR PRINTS: PR00001; GLABLOOD.	
DR SMART: SM00179; EGF_CA; 1.	
DR SMART: SM00001; EGF-like; 1.	
DR SMART: SM00069; GLA; 1.	
DR SMART: SM00020; TRYP_Spc; 1.	
DR PROSITE: PS00010; ASX_HYDROXYL; UNKNOWN_1.	
DR PROSITE: PS00022; EGF_1; UNKNOWN_1.	
DR PROSITE: PS01186; EGF_2; 2.	
DR PROSITE: PS01187; EGF_CA; 1.	
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.	
DR PROSITE: PS02040; TRYPSIN_DOM; 1.	
DR PROSITE: PS00134; TRYPSIN_HIS; UNKNOWN_1.	
DR PROSITE: PS00135; TRYPSIN_SER; 1.	
KW Calcium-binding; EGF-like domain; Glycoprotein; Hydrolase; Repeat;	
KM Serine protease; Signal.	
FT SIGNAL	POTENTIAL.
FT CHAIN	COAGULATION FACTOR X.
SQ SEQUENCE	481 AA: 54018 MM; 8AC09DE5EF9D271E CRC64;
Query Match	49.2%; Score 98; DB 11; Length 481;
Best Local Similarity	36.4%; Pred. No. 2e-08;
Matches	16; Conservative 10; Mismatches 16; Indels 0; Gaps 0;
QY	1 ANSFLXLRGSLXRXICIXXICDFFXAKXIFZDVDDTLAFWSKH 44
DB	41 ANSFEFFKKGNLRECMELCSYEFVRLEFDEDEKTKYWKY 84
RESULT 8	
Q96P08	PRELIMINARY; PRT: 701 AA.
AC Q96P08:	
DT 01-DEC-2001 (TREMblrel. 19, Created)	
DT 01-DEC-2001 (TREMblrel. 19, Last sequence update)	
DT 01-MAR-2002 (TREMblrel. 20, Last annotation update)	
OS Factor VII active site mutant immunocoujugate.	
OS Homo sapiens (Human).	
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.	
OC NCBI_TaxID=9606;	
RN [1]	
RP SEQUENCE FROM N.A.	
RX MEDLINE=21477448; PubMed=11593034;	
RA Hu Z., Garen A.;	
RT "Targeting tissue factor on tumor vascular endothelial cells and tumor	
RT cells for immunotherapy in mouse models of prostatic cancer.";	
RL Proc. Natl. Acad. Sci. U.S.A. 98:12180-12185(2001).	
DR EMBL: AF272774; AKS5686.1; "	
DR InterPro: IPR000152; Asx_hydroxyl.	
DR InterPro: IPR000561; EGF-like.	
DR InterPro: IPR000742; EGF_2.	
DR InterPro: IPR001881; EGF_Ca.	
DR InterPro: IPR003006; I9_MHC.	
DR InterPro: IPR001254; Ser_protease_Try.	
DR InterPro: IPR000294; Vltk_dep_GLA	

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DR Pfam: PF00008; EGF; 2.
DR Pfam: PF00594; gla; 1.
DR Pfam: PF00047; 1g; 2.
DR Pfam: PF00089; trypsin; 1.
DR SMART: SM00181; EGF; 2.
DR PROSITE: PS00010; ASX_HYDROXYL; UNKNOWN_1.
DR PROSITE: PS00022; EGF_1; UNKNOWN_1.
DR PROSITE: PS01186; EGF_2; UNKNOWN_1.
DR PROSITE: PS01187; EGF_CA; UNKNOWN_1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; UNKNOWN_1.
DR PROSITE: PS00290; IG_MHC; UNKNOWN_1.
DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; UNKNOWN_1.
DR PROSITE: PS00135; TRYPSIN_SER; UNKNOWN_1.
DR Hydrolase: Serine protease.
SQ SEQUENCE 701 AA; 77826 MW; 94AC6CEB42CC992F CRC64;

Query Match 49.2%; Score 98; DB 4; Length 701;
Best Local Similarity 48.8%; Pred. No. 3.1e-08;
Matches 20; Conservative 4; Mismatches 17; Indels 0; Gaps 0;

QY 1 ANSFLXLRHGSIXRXCIXICDPXXAKXIFZVDVDTLAFW 41
DB 61 ANAFLEELRPGSLERCKECCSFEBARLTFKDAERTKLEW 101

RESULT 9
Q9GMD9 PRELIMINARY; PRT; 469 AA.
ID Q9GMD9;
AC Q9GMD9;
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)
DE Coagulation factor X.
OS Ornithorhynchus anatinus (Duckbill platypus).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Monotremata; Ornithorhynchidae; Ornithorhynchus.
NCBI_TaxID=9258;
RN (1)
RP MEDLINE=21015017; PubMed=11132153;
RX "Poorafshar M., Aveshogh M., Kunday B., Hellman L.;
RT "Identification and structural analysis of four serine proteases in a
RT monotreme, the platypus, Ornithorhynchus anatinus."
CC Immunogenetics 52:19-28(2000).
CL -1 SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
CC TRYPSIN FAMILY
DR EMBL: AF215654; AAC00453.1; -.
DR HSSP: P00742; 1XKB.
DR MEROPS: S01.216; -.
DR InterPro: IPR000152; Asx_hydroxyl.
DR InterPro: IPR001314; Chymotrypsin.
DR InterPro: IPR000561; EGF-like.
DR InterPro: IPR000742; EGF_2.
DR InterPro: IPR001881; EGF_CA.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Ser_protease_Try.
DR InterPro: IPR000294; Vltk_dep_GLA.
DR Pfam: PF00008; EGF; 2.
DR Pfam: PF00594; gla; 1.
DR PRINTS: PR00089; trypsin; 1.
DR PRINTS: PR00722; CHYMOTRYPSIN.
DR SMART: SM00001; EGF; 2.
DR SMART: SM00179; EGF_CA; 1.
DR SMART: SM00001; EGF-like; 2.
DR SMART: SM00069; GLA; 1.
DR SMART: SM00020; TRYPSIN_SPEC; 1.
DR PROSITE: PS00010; ASX_HYDROXYL; UNKNOWN_1.
DR PROSITE: PS00022; EGF_1; UNKNOWN_1.
DR PROSITE: PS01186; EGF_2; 2.
DR PROSITE: PS01187; EGF_CA; 1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.

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DR PROSITE: PS00240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_HIS; UNKNOWN_1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
DR Hydrolase: Serine protease.
SQ SEQUENCE 469 AA; 52196 MW; 4C66C230D0758F6A CRC64;

Query Match 46.2%; Score 92; DB 6; Length 469;
Best Local Similarity 38.1%; Pred. No. 2.2e-07;
Matches 16; Conservative 7; Mismatches 19; Indels 0; Gaps 0;

QY 1 ANSFLXLRHGSIXRXCIXICDPXXAKXIFZVDVDTLAFWS 42
DB 41 ANSLEELKGNLRECNETCSYEAREVFEDTQTNFEWN 82

RESULT 10
Q15253 PRELIMINARY; PRT; 100 AA.
ID Q15253;
AC Q15253;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE Thrombin precursor (fragment).
GN F2.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
NCBI_TaxID=9606;
RN (1)
RP SEQUENCE FROM N.A.
RX MEDLINE=87182874; PubMed=3471151;
RA Macgillivray R.T., Irwin D.M., Guinto E.R., Stone J.C.;
RT "Recombinant genetic approaches to functional mapping of thrombin."
RL Ann. N.Y. Acad. Sci. 485:73-79(1986).
DR EMBL: M3031; AAC60220.1; -.
DR HSSP: P00735; 2PFI.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR000294; Vltk_dep_GLA.
DR Pfam: PF00594; gla; 1.
DR PRINTS: PR00001; GLABLOOD.
DR SMART: SM00069; GLA; 1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR SIGNAL.
FT SIGNAL. 1 43 POTENTIAL.
FT CHAIN 44 >100 POTENTIAL.
FT NON_TER 100 100
SQ SEQUENCE 100 AA; 11302 MW; FDE05D0174E1F6FE CRC64;

Query Match 40.7%; Score 81; DB 4; Length 100;
Best Local Similarity 34.1%; Pred. No. 3.3e-06;
Matches 15; Conservative 8; Mismatches 21; Indels 0; Gaps 0;

QY 1 ANSFLXLRHGSIXRXCIXICDPXXAKXIFZVDVDTLAFWSKH 44
DB 44 ANTFLEEVKGNLRECNETCSYEAREVFLESTATDVFMKY 87

RESULT 11
Q61109 PRELIMINARY; PRT; 446 AA.
ID Q61109;
AC Q61109;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)
DE Coagulation factor VII.
GN F7 OR FVII.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NCBI_TaxID=10090;
RN (1)
RP SEQUENCE FROM N.A.
RC TISSUE=LIVER;

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RX MEDLINE=96276538; PubMed=8701412;
RA Idusogie E., Rosen E., Geng J.P., Carmeliet P., Collen D.,
RA Castellino F.J.;
RT "Characterization of a cDNA encoding murine coagulation factor VII.";
RT Thromb. Haemost. 75:481-487(1996).
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
CC TRYPSIN FAMILY.
DR EMBL: U44795; AAC52570.1; -.
DR HSSP: P08709; 1FAK.
DR MEROPS: S01.215; -.
DR MGD: MGI:109325; F7.
DR InterPro: IPR002086; Aldehyde_dehydr.
DR InterPro: IPR000152; ASX_hydroxyl.
DR InterPro: IPR001314; Chymotrypsin.
DR InterPro: IPR01064; Crystallin.
DR InterPro: IPR000561; EGF-like.
DR InterPro: IPR001881; EGF_Ca.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001254; Ser_protease_Try.
DR InterPro: IPR00294; VitK_dep_GLA.
DR Pfam: PF00008; EGF_2.
DR Pfam: PF00594; gla; 1.
DR Pfam: PF00089; trypsin; 1.
DR PRINTS: PRO0722; CHYMOTRYPSIN.
DR PRINTS: PRO0001; GLABLOOD.
DR SMART: SM00179; EGF_CA; 1.
DR SMART: SM00001; EGF-like; 1.
DR SMART: SM00069; GLA; 1.
DR SMART: SM00020; TRYP_SPC; 1.
DR PROSITE: PS00070; ALDEHYDE_DEHYDR_CYS; UNKNOWN_1.
DR PROSITE: PS00010; ASX_HYDROXYL; UNKNOWN_1.
DR PROSITE: PS00225; CRYSTALLIN_BETAGAMMA; UNKNOWN_1.
DR PROSITE: PS00022; EGF_1; UNKNOWN_1.
DR PROSITE: PS01187; EGF_CA; 1.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR PROSITE: PS50240; TRYPSIN_DOM; 1.
DR PROSITE: PS00134; TRYPSIN_SIS; UNKNOWN_1.
DR PROSITE: PS00135; TRYPSIN_SER; 1.
DR Calcium-binding; EGF-like domain; Glycoprotein; Hydrolase; Repeat;
KW Serine protease.
SQ SEQUENCE 446 AA; 50318 MW; 482FD09BEFDA6870 CRC64;

Query Match 39.7%; Score 79; DB 11; Length 446;
Best Local Similarity 43.9%; Pred. No. 3.8e-05;
Matches 18; Conservative 3; Mismatches 20; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSILXRXCIXICDFXAKXIFZVDVDTLAFW 41
DB 42 ANSLEELMPGSLERECNEQCSFEARERIFKSPERTKQFW 82

RESULT 12
OQNSDO PRELIMINARY: PRT: 650 AA.
AC OQNSDO:
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)
DE Protein S precursor.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OC NCBI_Taxid=9606;
OX [1]
RN SEQUENCE FROM N.A.
RC TISSUE=LIVER;
RA Wydro R., Cohen E., Dackowski W., Stenflo J., Lundwall A.,
RA Dahlback B.;
RL Submitted (FEB-1992) to the EMBL/Genbank/DBJ databases.
DR EMBL: X12892; CAA31383.1; -.
DR HSSP: P00740; 1CFH.
DR InterPro: IPR000152; ASX_hydroxyl.
DR InterPro: IPR000561; EGF-like.

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DR InterPro: IPR001881; EGF_Ca.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001791; Laminin_G.
DR InterPro: IPR00294; VitK_dep_GLA.
DR Pfam: PF00008; EGF_4.
DR Pfam: PF00594; gla; 1.
DR Pfam: PF00054; laminin_G; 1.
DR PRINTS: PRO0001; GLABLOOD.
DR SMART: SM00179; EGF_CA; 3.
DR SMART: SM00069; GLA; 1.
DR SMART: SM00282; Lamg; 2.
DR PROSITE: PS00010; ASX_HYDROXYL; 3.
DR PROSITE: PS00022; EGF_1; UNKNOWN_1.
DR PROSITE: PS01186; EGF_2; 3.
DR PROSITE: PS01187; EGF_CA; 2.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR Calcium-binding; EGF-like domain; Glycoprotein; Hydroxylation; Repeat;
KW signal.
FT SIGNAL 1 15 POTENTIAL.
FT CHAIN 16 650 POTENTIAL.
SQ SEQUENCE 650 AA; 72480 MW; C67345CEB645174 CRC64;

Query Match 39.7%; Score 79; DB 4; Length 650;
Best Local Similarity 34.1%; Pred. No. 5.7e-05;
Matches 15; Conservative 11; Mismatches 18; Indels 0; Gaps 0;

OY 1 ANSFLXLRHGSILXRXCIXICDFXAKXIFZVDVDTLAFWSKH 44
DB 16 ANSLEETKQGLERECIEELCNKEAREVFNDETFYFKY 59

RESULT 13
OQ16519 PRELIMINARY: PRT: 650 AA.
AC OQ16519:
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE Protein S precursor (Fragment).
GN Prosl.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OC NCBI_Taxid=9606;
OX [1]
RN SEQUENCE FROM N.A.
RX MEDLINE=86313649; PubMed=2944113;
RA Lundwall A., Dackowski W., Cohen E., Shaffer M., Mahr A., Dahlback B.,
RA Stenflo J., Wydro R.;
RT "Isolation and sequence of the cDNA for human protein S, a regulator
RT of blood coagulation.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:6716-6720(1986).
DR EMBL: M14338; AAA60181.1; -.
DR HSSP: P00740; 1CFH.
DR InterPro: IPR000152; ASX_hydroxyl.
DR InterPro: IPR000561; EGF-like.
DR InterPro: IPR001881; EGF_Ca.
DR InterPro: IPR002383; GLA_blood.
DR InterPro: IPR001791; Laminin_G.
DR InterPro: IPR000294; VitK_dep_GLA.
DR Pfam: PF00008; EGF_4.
DR Pfam: PF00594; gla; 1.
DR Pfam: PF00054; laminin_G; 1.
DR PRINTS: PRO0001; GLABLOOD.
DR SMART: SM00179; EGF_CA; 3.
DR SMART: SM00069; GLA; 1.
DR SMART: SM00282; Lamg; 2.
DR PROSITE: PS00010; ASX_HYDROXYL; 3.
DR PROSITE: PS00022; EGF_1; UNKNOWN_1.
DR PROSITE: PS01186; EGF_2; 3.
DR PROSITE: PS01187; EGF_CA; 2.
DR PROSITE: PS00011; GLU_CARBOXYLATION; 1.
DR Calcium-binding; EGF-like domain; Glycoprotein; Hydroxylation; Repeat;
KW

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SEQUENCE				
Query Match		39.7%;	Score 79;	DB 4; Length 650;
Best Local Similarity		34.1%;	Pred. NO. 5.7e-05;	
Matches	15;	Conservative	11;	Mismatches 18; Indels 0; Gaps 0
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DB	16	ANSLEETKQGNLERCEIEELCKNEAREAREVFNDEPDTDFYKX	59	
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AC	O8T6I3:			
DT	01-JUN-2002 (TREMBLrel. 21, Created)			
DT	01-JUN-2002 (TREMBLrel. 21, Last sequence update)			
DT	01-JUN-2002 (TREMBLrel. 21, Last annotation update)			
DE	Gla-like protein.			
OS	Halocynthia roretzi (Sea squirt).			
OC	Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea;			
CC	Stolidobranchia; Pyuridae; Halocynthia.			
OX	NCBI_TaxID=7729;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RT	Wang C.-P., Stafford D.W.;			
RL	"Halocynthia roretzi gla-like protein partial genomic DNA sequence."			
DR	Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.			
DR	EMBL: AF466701; AAL74247.2; "			
SEQUENCE	542 AA;	62090 MW;	EB9BF13FE42B32FE	CRC64;
Query Match		38.4%;	Score 76.5;	DB 5; Length 542;
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DB	33	SHEEIOGNGLERCEIEELCSFEAREAREVFNQIDNERNARY	75	
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O28994				
ID	O28994	PRELIMINARY;	PRT;	138 AA.
AC	O28994:			
DT	01-NOV-1996 (TREMBLrel. 01, Created)			
DT	01-NOV-1996 (TREMBLrel. 01, Last sequence update)			
DT	01-DEC-2001 (TREMBLrel. 19, Last annotation update)			
DE	Mature porcine factor IX (Fragment).			
OS	Sus scrofa (pig).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
CC	Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.			
OX	NCBI_TaxID=9823;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RT	TISSUE=LIVER;			
RL	MEDLINE=96003866; PubMed=7568220;			
RA	Brandstetter H., Bauer M., Huber R., Lollar P., Bode W.;			
RT	"X-ray structure of clotting factor IXa: active site and module			
PROC.	structure related to zase activity and hemophilia B."			
PROC.	Natl. Acad. Sci. U.S.A. 92:9796-9800(1995).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RT	TISSUE=LIVER;			
RL	Lollar P.;			
DR	Submitted (MAR-1996) to the EMBL/GenBank/DBJ databases.			
DR	EMBL: U51135; AAA96318.1; "			
DR	HSSP: P00740; 1EDM.			
DR	InterPro: IPR000152; Asx_hydroxyl.			
DR	InterPro: IPR000561; EGF-like.			

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DR      InterPro: IPR000742; EGF_2.
DR      InterPro: IPR001881; EGF_CA.
DR      InterPro: IPR001438; EGF_IT.
DR      InterPro: IPR002383; GLA_blood.
DR      InterPro: IPR000294; VltK_dep_GLA.
DR      Pfam: PF000068; EGF_2.
DR      Pfam: PF00594; gla; 1.
DR      PRINTS: PR000010; EGFBLDOD.
DR      PRINTS: PR00001; GLABLOD.
DR      SMART: SM00179; EGF_CA; 1.
DR      SMART: SM00069; GLA; 1.
DR      PROSITE: PS00010; ASX_HYDROXYL; UNKNOWN_1.
DR      PROSITE: PS00022; EGF_1; UNKNOWN_1.
DR      PROSITE: PS01186; EGF_2; 2.
DR      PROSITE: PS01187; EGF_CA; 1.
KW      Calcium-binding; EGF-like domain; Glycoprotein; Repeat.
FT      NON_TER          1
FT      FT              138
SQ      SEQUENCE        138 AA; 15515 MW; 793BABADEAD5FAFD CXC64;
Query Match           38.2%; Score 76; DB 6; Length 136;
Best Local Similarity 35.3%; Pred. No. 3.4e-05;
Matches 12; Conservative 8; Mismatches 14; Indels 0; Gaps 0;
QY    11 GSLSRXCIXICDFXXAKKIETZVDVDTIAFWSKH 44
       ||||| | | : : : : | | : :
DB    4 GNLERECIEKCSFEAREVFENTERTENFWKY 37

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Search completed: May 13, 2003, 14:04:10  
Job time : 31 secs

FW	Signal.	1	1	
FT	NON_TER	<1	15	POTENTIAL.
FT	SIGNAL	16	650	PROTEIN S.
SO	CHAIN	650 AA:	72462 MW;	9A8C044C503BF474 CRC64;
SEQUENCE				
Query Match		39.7%;	Score 79;	DB 4; Length 650;
Best Local Similarity		34.1%;	Pred. NO. 5.7e-05;	
Matches	15;	Conservative	11;	Mismatches 18; Indels 0; Gaps 0
OY	1	ANSFLXLRHGSGLXRCXCIXXICDPFXAKXIFZDVDDTLAFWSKH	44	
DB	16	ANSLEETKQGNLERCEIEELCKNEAREAREVFNDEPDTDFYKX	59	
RESULT 14				
O8T6I3				
ID	O8T6I3	PRELIMINARY;	PRT;	542 AA.
AC	O8T6I3:			
DT	01-JUN-2002 (TREMBLrel. 21, Created)			
DT	01-JUN-2002 (TREMBLrel. 21, Last sequence update)			
DT	01-JUN-2002 (TREMBLrel. 21, Last annotation update)			
DE	Gla-like protein.			
OS	Halocynthia roretzi (Sea squirt).			
OC	Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea;			
CC	Stolidobranchia; Pyuridae; Halocynthia.			
OX	NCBI_TaxID=7729;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RT	Wang C.-P., Stafford D.W.;			
RL	"Halocynthia roretzi gla-like protein partial genomic DNA sequence."			
DR	Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.			
DR	EMBL: AF466701; AAL74247.2; "			
SEQUENCE	542 AA;	62090 MW;	EB9BF13FE42B32FE	CRC64;
Query Match		38.4%;	Score 76.5;	DB 5; Length 542;
Best Local Similarity		30.2%;	Pred. NO. 0.00013;	
Matches	13;	Conservative	11;	Mismatches 18; Indels 1; Gaps 1
OY	3	SFLXLRHGSGLXRCXCIXXICDPFXAKXIFZ-DVDDTLAFWSKH	44	
DB	33	SHEEIOGNGLERCEIEELCSFEAREAREVFNQIDNERNARY	75	
RESULT 15				
O28994				
ID	O28994	PRELIMINARY;	PRT;	138 AA.
AC	O28994:			
DT	01-NOV-1996 (TREMBLrel. 01, Created)			
DT	01-NOV-1996 (TREMBLrel. 01, Last sequence update)			
DT	01-DEC-2001 (TREMBLrel. 19, Last annotation update)			
DE	Mature porcine factor IX (Fragment).			
OS	Sus scrofa (pig).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
CC	Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.			
OX	NCBI_TaxID=9823;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RT	TISSUE=LIVER;			
RL	MEDLINE=96003866; PubMed=7568220;			
RA	Brandstetter H., Bauer M., Huber R., Lollar P., Bode W.;			
RT	"X-ray structure of clotting factor IXa: active site and module			
PROC.	structure related to zase activity and hemophilia B."			
PROC.	Natl. Acad. Sci. U.S.A. 92:9796-9800(1995).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RT	TISSUE=LIVER;			
RL	Lollar P.;			
DR	Submitted (MAR-1996) to the EMBL/GenBank/DBJ databases.			
DR	EMBL: U51135; AAA96318.1; "			
DR	HSSP: P00740; 1EDM.			
DR	InterPro: IPR000152; Asx_hydroxyl.			
DR	InterPro: IPR000561; EGF-like.			

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## OM protein - protein search, using sw model

Run on: May 13, 2003, 14:03:19 ; Search time 15 Seconds

(without alignments)  
86.307 Million cell updates/sec

Title: SEQ1-ED1RED

Perfect score: 199

Sequence: 1 ANSEFLXLRHSGSLXRCXCIX.....XXAKXIFZVDDTLAFWSKH 44

Scoring table: BIOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 262574 seqs, 29422922 residues

Total number of hits satisfying chosen parameters: 262574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

Issued\_Patents\_AA:\*  
1: /cgn2\_6/ptodata/1/1aa/5A.COMB.pep:\*  
2: /cgn2\_6/ptodata/1/1aa/5B.COMB.pep:\*  
3: /cgn2\_6/ptodata/1/1aa/6A.COMB.pep:\*  
4: /cgn2\_6/ptodata/1/1aa/6B.COMB.pep:\*  
5: /cgn2\_6/ptodata/1/1aa/PCVUS.COMB.pep:\*  
6: /cgn2\_6/ptodata/1/1aa/backfileseq1.pep:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	175	87.9	44	3	US-08-955-636-19
2	173	86.9	44	3	US-08-955-636-24
3	173	86.9	44	3	US-08-955-636-35
4	169	84.9	44	3	US-08-955-636-1
5	169	84.9	45	2	US-08-955-832-2
6	169	84.9	419	1	US-08-295-411-1
7	169	84.9	419	2	US-08-955-471-1
8	169	84.9	419	4	US-09-667-570A-3
9	169	84.9	419	5	PCR-US92-10242-1
10	169	84.9	460	2	US-08-756-506-2
11	169	84.9	460	6	5270178-13
12	169	84.9	460	6	5270178-14
13	169	84.9	460	6	5270178-15
14	169	84.9	460	6	5270178-16
15	169	84.9	461	6	5225537-2
16	169	84.9	461	6	5270178-17
17	169	84.9	461	6	5270178-18
18	169	84.9	461	6	5270178-19
19	169	84.9	461	6	5460953-3
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21	167	83.9	44	3	US-08-955-636-21
22	167	83.9	44	3	US-08-955-636-25
23	166	83.4	44	3	US-08-955-636-22
24	156	78.4	42	2	US-08-745-254A-2
25	156	78.4	461	6	5270178-2
26	152	76.4	41	1	US-08-229-280-5
27	138	69.3	410	4	US-09-065-872-1

28	138	69.3	410	4	US-09-667-570A-1	Sequence 1, Appl
29	130	65.3	409	4	US-09-065-872-2	Sequence 2, Appl
30	130	65.3	409	4	US-09-667-570A-2	Sequence 2, Appl
31	126	63.3	44	3	US-08-955-636-23	Sequence 23, Appl
32	116	58.3	44	3	US-08-955-636-2	Sequence 2, Appl
33	112	56.3	139	1	US-08-330-978-2	Sequence 2, Appl
34	112	56.3	139	1	US-08-474-042-2	Sequence 2, Appl
35	112	56.3	139	1	US-08-484-558-2	Sequence 2, Appl
36	112	56.3	139	1	US-08-774-593-2	Sequence 2, Appl
37	112	56.3	437	1	US-08-487-037-2	Sequence 2, Appl
38	112	56.3	437	1	US-08-487-037-3	Sequence 3, Appl
39	112	56.3	488	1	US-08-487-037-1	Sequence 3, Appl
40	108	54.3	487	1	US-08-469-486-53	Sequence 53, Appl
41	108	54.3	487	2	US-08-469-658-53	Sequence 2, Appl
42	108	54.3	492	2	US-08-469-486-2	Sequence 2, Appl
43	108	54.3	492	2	US-08-469-658-2	Sequence 2, Appl
44	107	53.8	448	1	US-08-295-411-3	Sequence 3, Appl
45	107	53.8	448	2	US-08-955-471-3	Sequence 3, Appl

## ALIGNMENTS

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RESULT 1
US-08-955-636-19
; Sequence 19, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelstuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; NUMBER OF SEQ ID NOS: 35
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 19
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: MOD_RES
; LOCATION: (0)...(0)
; OTHER INFORMATION: Xaa-gamma carboxyglutamic acid or glutamic acid
US-08-955-636-19

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Best Local Similarity 95.5%; Pred. No. 1.2e-22;
Matches 42; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY      1 ANSEFLXLRHSGSLXRCXCIXICDFFXAKXIFZVDDTLAFWSKH 44
Db      1 ANSEFLXLRHSGSLXRCXCIXICDFFXAKXIFZVDDTLAFWSKH 44

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US-08-955-636-24
; Sequence 24, Application US/08955636A
; Patent No. 6017882
; GENERAL INFORMATION:
; APPLICANT: Nelstuen, Gary
; TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT
; FILE REFERENCE: 09531/002001
; CURRENT APPLICATION NUMBER: US/08/955,636A
; CURRENT FILING DATE: 1997-10-23
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 24
; LENGTH: 44
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:

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NAME/KEY: MOD\_RES  
 LOCATION: (0)...(0)  
 OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid  
 US-08-955-636-24

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RESULT 3  
 US-08-955-636-35

Sequence 35, Application US/08955636A

Patent No. 6017862

GENERAL INFORMATION:

APPLICANT: Nelsestuen, Gary

TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT

TITLE OF INVENTION: POLYPEPTIDES

FILE REFERENCE: 09531/002001

CURRENT APPLICATION NUMBER: US/08/955,636A

CURRENT FILING DATE: 1997-10-23

NUMBER OF SEQ ID NOS: 35

SOFTWARE: FastSeq for Windows Version 3.0

SEQ ID NO: 35

LENGTH: 44

TYPE: PRT

ORGANISM: Homo sapiens

FEATURE:

NAME/KEY: MOD\_RES

LOCATION: (0)...(0)

OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid

US-08-955-636-35

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 DB 1 ANSFLXLRHGSILRXICIXICDFFXAKXIFZVDVDTLAFWSKH 44

RESULT 4  
 US-08-955-636-1

Sequence 1, Application US/08955636A

Patent No. 6017862

GENERAL INFORMATION:

APPLICANT: Nelsestuen, Gary

TITLE OF INVENTION: MODIFIED VITAMIN K-DEPENDENT

TITLE OF INVENTION: POLYPEPTIDES

FILE REFERENCE: 09531/002001

CURRENT APPLICATION NUMBER: US/08/955,636A

CURRENT FILING DATE: 1997-10-23

NUMBER OF SEQ ID NOS: 35

SOFTWARE: FastSeq for Windows Version 3.0

SEQ ID NO: 1

LENGTH: 44

TYPE: PRT

ORGANISM: Homo sapiens

FEATURE:

NAME/KEY: MOD\_RES

LOCATION: (0)...(0)

OTHER INFORMATION: Xaa=gamma carboxyglutamic acid or glutamic acid

US-08-955-636-1

Query Match 84.9%; Score 169; DB 3; Length 44;  
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RESULT 5  
 US-08-965-832-2

Sequence 2, Application US/08965832

Patent No. 5847085

GENERAL INFORMATION:

APPLICANT: CHARLES T. ESMON AND MIKHAIL D. SMIRNOV

TITLE OF INVENTION: Modified Protein C

NUMBER OF SEQUENCES: 4

CORRESPONDENCE ADDRESS:

ADDRESSEE: Patricia L. Pabst

STREET: 2800 One Atlantic Center, 1201 West

STREET: Peachtree Street

CITY: Atlanta

STATE: GA

COUNTRY: USA

ZIP: 30309-3450

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/965,832

FILING DATE: 7-NOV-1997

CLASSIFICATION: 530

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/745,254

FILING DATE: 8-NOV-1996

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 60/053,768

FILING DATE: 25-JUL-1997

ATTORNEY/AGENT INFORMATION:

NAME: Pabst, Patricia L.

REGISTRATION NUMBER: 31,284

REFERENCE/DOCKET NUMBER: OMRF 165/167

TELECOMMUNICATION INFORMATION:

TELEPHONE: (404)-873-8794

TELEFAX: (404)-873-8795

INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:

LENGTH: 45 amino acids

TYPE: amino acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: peptide

FEATURE:

NAME/KEY:

LOCATION: 6, 7, 14, 16, 19, 20, 25, 26, 29

OTHER INFORMATION: /note="where Xaa means gamma

OTHER INFORMATION: carboxyglutamic acid"

FEATURE:

NAME/KEY:

LOCATION:

OTHER INFORMATION: /note="partial sequence of human protein C"

US-08-965-832-2

Query Match 84.9%; Score 169; DB 2; Length 45;  
 Best Local Similarity 93.2%; Pred. No. 1.2e-21;  
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 DB 1 ANSFLXLRHGSILRXICIXICDFFXAKXIFZVDVDTLAFWSKH 44

RESULT 6  
 US-08-295-411-1  
 Sequence 1, Application US/08295411

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Patent No. 5679639
GENERAL INFORMATION:
APPLICANT: Griffin, John H.
APPLICANT: Masters, Rolf M.
TITLE OF INVENTION: Serine Protease-Derived Polypeptides and
TITLE OF INVENTION: Anti-Peptide Antibodies, Systems and Therapeutic Methods
TITLE OF INVENTION: for Inhibiting Coagulation
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESS: Office of Patent Counsel, The Scripps
ADDRESS: Research Institute
STREET: 10666 No. 5679639th Torrey Pines Road, TPC 8
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/295,411
FILING DATE: 22-AUG-1994
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/793,989
FILING DATE: 18-NOV-1991
CLASSIFICATION: 530
ATTORNEY/AGENT INFORMATION:
NAME: Filling, Thomas
REGISTRATION NUMBER: 34,163
REFERENCE/DOCKET NUMBER: TSRI263.0C1
TELEPHONE: 619-554-2937
TELEFAX: 619-554-6312
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 419 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: Region
LOCATION: 1..157
OTHER INFORMATION: /note="Protein C Light Chain"
FEATURE:
NAME/KEY: Region
LOCATION: 158..169
OTHER INFORMATION: /note="Protein C Activation
Peptide"
FEATURE:
NAME/KEY: Region
LOCATION: 170..419
OTHER INFORMATION: /note="Protein C Heavy Chain"
US-08-295-411-1
Query Match 84.9%; Score 169; DB 1; Length 419;
Best Local Similarity 72.7%; Pred. No. 1.5e-20;
Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;
QY 1 ANSFLLXLRHGSIXRCIXIXICDFFXAXAKIFZVDVDTLAFWSKH 44
DB 1 ANSFLELRHSSLERCIEICDFFXAXAKIFZVDVDTLAFWSKH 44
RESULT 7
US-08-955-471-1
Sequence 1, Application US/08955471
Patent No. 5968751
GENERAL INFORMATION:
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APPLICANT: Griffin, John H.
APPLICANT: Masters, Rolf M.
TITLE OF INVENTION: Serine Protease-Derived Polypeptides and
TITLE OF INVENTION: Anti-Peptide Antibodies, Systems and Therapeutic Methods
TITLE OF INVENTION: for Inhibiting Coagulation
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESS: Office of Patent Counsel, The Scripps
ADDRESS: Research Institute
STREET: 10666 No. 5968751th Torrey Pines Road, TPC 8
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/955,471
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/295,411
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Filling, Thomas
REGISTRATION NUMBER: 34,163
REFERENCE/DOCKET NUMBER: TSRI263.0C1
TELEPHONE: 619-554-2937
TELEFAX: 619-554-6312
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 419 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
HYPOTHETICAL: NO
ANTI-SENSE: NO
FEATURE:
NAME/KEY: Region
LOCATION: 1..157
OTHER INFORMATION: /note="Protein C Light Chain"
FEATURE:
NAME/KEY: Region
LOCATION: 158..169
OTHER INFORMATION: /note="Protein C Activation
Peptide"
FEATURE:
NAME/KEY: Region
LOCATION: 170..419
OTHER INFORMATION: /note="Protein C Heavy Chain"
US-08-955-471-1
Query Match 84.9%; Score 169; DB 2; Length 419;
Best Local Similarity 72.7%; Pred. No. 1.5e-20;
Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;
QY 1 ANSFLLXLRHGSIXRCIXIXICDFFXAXAKIFZVDVDTLAFWSKH 44
DB 1 ANSFLELRHSSLERCIEICDFFXAXAKIFZVDVDTLAFWSKH 44
RESULT 8
US-09-667-570A-3
Sequence 3, Application US/09667570A
Patent No. 6436397
GENERAL INFORMATION:
APPLICANT: Baker, Jeffrey C
APPLICANT: Carlson, Andrew D
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; APPLICANT: Huang, Lihua
; APPLICANT: Shelliga, Theodore A
; TITLE OF INVENTION: Improved Methods for Processing Activated Protein C
; FILE REFERENCE: X-11796A
; CURRENT APPLICATION NUMBER: US/09/667,570A
; CURRENT FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: 60/045,255
; PRIOR FILING DATE: 1997-04-28
; NUMBER OF SEQ ID NOS: 3
; SOFTWARE: Patentin version 3.1
; SEQ ID NO: 3
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
; US-09-667-570A-3

Query Match
Best Local Similarity 84.9%; Score 169; DB 4; Length 419;
Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;

OY 1 ANSFLXXLRHGSILRXICIXXICDFFXXAKXIFZDVDDTLAFWSKH 44
Db 1 ANSFLRLRHSLSLRECIETICDFEAKELFQVNDTLAFWSKH 44

RESULT 9
PCT-US92-10242-1
; Sequence 1, Application PC/TUS9210242
; GENERAL INFORMATION:
; APPLICANT: Griffin, John H.
; APPLICANT: Masters, Rolf
; TITLE OF INVENTION: Serine Protease-Derived Polypeptides and
; TITLE OF INVENTION: Anti-Peptide Antibodies, Systems and Therapeutic Methods
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Office of Patent Counsel, The Scripps
; ADDRESSEE: Research Institute
; STREET: 10666 North Torrey Pines Road, TPC 8
; CITY: La Jolla
; STATE: CA
; COUNTRY: USA
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/10242
; FILING DATE: 19921118
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/793,989
; FILING DATE: 18-NOV-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Fitting, Thomas
; REGISTRATION NUMBER: 34,163
; REFERENCE/DOCKET NUMBER: SCRO472P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619-554-2937
; TELEFAX: 619-554-6312
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 419 amino acids
; TYPE: AMINO ACID
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; FEATURE:
; NAME/KEY: Region
; LOCATION: 1..157
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; OTHER INFORMATION: /note="Protein C Light Chain"
; FEATURE:
; NAME/KEY: Region
; LOCATION: 158..169
; OTHER INFORMATION: /note="Protein C Activation"
; OTHER INFORMATION: Peptide"
; FEATURE:
; NAME/KEY: Region
; LOCATION: 170..419
; OTHER INFORMATION: /note="Protein C Heavy Chain"
; PCT-US92-10242-1

Query Match
Best Local Similarity 84.9%; Score 169; DB 5; Length 419;
Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;

OY 1 ANSFLXXLRHGSILRXICIXXICDFFXXAKXIFZDVDDTLAFWSKH 44
Db 1 ANSFLRLRHSLSLRECIETICDFEAKELFQVNDTLAFWSKH 44

RESULT 10
US-08-756-506-2
; Sequence 2, Application US/08756506
; Patent No. 5905185
; GENERAL INFORMATION:
; APPLICANT: Garner, Ian
; APPLICANT: Cottingham, Ian R.
; APPLICANT: Temperley, Simon M.
; APPLICANT: Foster, Donald C.
; APPLICANT: Sprecher, Cindy A.
; APPLICANT: Prunkard, Donna E.
; TITLE OF INVENTION: PROTEIN C PRODUCTION IN TRANSGENIC
; TITLE OF INVENTION: ANIMALS
; NUMBER OF SEQUENCES: 25
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Zymogenetics, Inc.
; STREET: 1201 Eastlake Avenue East
; CITY: Seattle
; STATE: WA
; COUNTRY: USA
; ZIP: 98102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/756,506
; FILING DATE:
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: Sawislak, Deborah A
; REGISTRATION NUMBER: 37,438
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 206-442-6672
; TELEFAX: 206-442-6672
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 460 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-756-506-2

Query Match
Best Local Similarity 84.9%; Score 169; DB 2; Length 460;
Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;

OY 1 ANSFLXXLRHGSILRXICIXXICDFFXXAKXIFZDVDDTLAFWSKH 44
Db 43 ANSFLRLRHSLSLRECIETICDFEAKELFQVNDTLAFWSKH 86
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RESULT 11  
US-08-756-506-4  
Sequence 4, Application US/08756506  
Patent No. 5905185  
GENERAL INFORMATION:  
APPLICANT: Garner, Ian  
APPLICANT: Cottingham, Ian R.  
APPLICANT: Temperley, Simon M.  
APPLICANT: Foster, Donald C.  
APPLICANT: Sprecher, Cindy A.  
APPLICANT: Prunkard, Donna E.  
TITLE OF INVENTION: PROTEIN C PRODUCTION IN TRANSGENIC  
NUMBER OF SEQUENCES: 25.  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: ZymoGenetics, Inc.  
STREET: 1201 Eastlake Avenue East  
CITY: Seattle  
STATE: WA  
COUNTRY: USA  
ZIP: 98102  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/756,506  
FILING DATE:  
CLASSIFICATION: 800  
ATTORNEY/AGENT INFORMATION:  
NAME: Sawislak, Deborah A  
REGISTRATION NUMBER: 37,438  
REFERENCE/DOCKET NUMBER: 95-28  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 206-442-6672  
TELEFAX: 206-442-6678  
INFORMATION FOR SEQ ID NO: 4:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 460 amino acids  
TYPE: amino acid  
TOPOLOGY: linear  
MOLECULE TYPE: protein  
US-08-756-506-4

Query Match 84.9%; Score 169; DB 2; Length 460;  
Best Local Similarity 72.7%; Pred. No. 1.7e-20;  
Matches 32: Conservative 2; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHGSGLXRXCIXXICDFXXAKXIFZVDVDTLAFWSKH 44  
Db 43 ANSFLLELRHSSLRECEIEICDFEAKEIFONVDDTLAFWSKH 86

RESULT 12  
5270178-13  
Patent No. 5270178  
APPLICANT: GERLITZ, BRUCE E.; GRINNELL, BRIAN W.  
TITLE OF INVENTION: VECTORS AND COMPOUNDS FOR EXPRESSION OF  
ZYMOGEN FORMS OF HUMAN PROTEIN C  
NUMBER OF SEQUENCES: 21  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/484,133  
FILING DATE: 23-FEB-1990  
SEQ ID NO: 13:  
LENGTH: 460  
5270178-13

Query Match 84.9%; Score 169; DB 6; Length 460;  
Best Local Similarity 72.7%; Pred. No. 1.7e-20;  
Matches 32: Conservative 2; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHGSGLXRXCIXXICDFXXAKXIFZVDVDTLAFWSKH 44  
Db 43 ANSFLLELRHSSLRECEIEICDFEAKEIFONVDDTLAFWSKH 86

RESULT 13  
5270178-14  
Patent No. 5270178  
APPLICANT: GERLITZ, BRUCE E.; GRINNELL, BRIAN W.  
TITLE OF INVENTION: VECTORS AND COMPOUNDS FOR EXPRESSION OF  
ZYMOGEN FORMS OF HUMAN PROTEIN C  
NUMBER OF SEQUENCES: 21  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/484,133  
FILING DATE: 23-FEB-1990  
SEQ ID NO: 14:  
LENGTH: 460  
5270178-14

Query Match 84.9%; Score 169; DB 6; Length 460;  
Best Local Similarity 72.7%; Pred. No. 1.7e-20;  
Matches 32: Conservative 2; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHGSGLXRXCIXXICDFXXAKXIFZVDVDTLAFWSKH 44  
Db 43 ANSFLLELRHSSLRECEIEICDFEAKEIFONVDDTLAFWSKH 86

RESULT 14  
5270178-15  
Patent No. 5270178  
APPLICANT: GERLITZ, BRUCE E.; GRINNELL, BRIAN W.  
TITLE OF INVENTION: VECTORS AND COMPOUNDS FOR EXPRESSION OF  
ZYMOGEN FORMS OF HUMAN PROTEIN C  
NUMBER OF SEQUENCES: 21  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/484,133  
FILING DATE: 23-FEB-1990  
SEQ ID NO: 15:  
LENGTH: 460  
5270178-15

Query Match 84.9%; Score 169; DB 6; Length 460;  
Best Local Similarity 72.7%; Pred. No. 1.7e-20;  
Matches 32: Conservative 2; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHGSGLXRXCIXXICDFXXAKXIFZVDVDTLAFWSKH 44  
Db 43 ANSFLLELRHSSLRECEIEICDFEAKEIFONVDDTLAFWSKH 86

RESULT 15  
5270178-16  
Patent No. 5270178  
APPLICANT: GERLITZ, BRUCE E.; GRINNELL, BRIAN W.  
TITLE OF INVENTION: VECTORS AND COMPOUNDS FOR EXPRESSION OF  
ZYMOGEN FORMS OF HUMAN PROTEIN C  
NUMBER OF SEQUENCES: 21  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/484,133  
FILING DATE: 23-FEB-1990  
SEQ ID NO: 16:  
LENGTH: 460  
5270178-16

Query Match 84.9%; Score 169; DB 6; Length 460;  
Best Local Similarity 72.7%; Pred. No. 1.7e-20;  
Matches 32: Conservative 2; Mismatches 10; Indels 0; Gaps 0;

Qy 1 ANSFLXLRHGSGLXRXCIXXICDFXXAKXIFZVDVDTLAFWSKH 44  
Db 43 ANSFLLELRHSSLRECEIEICDFEAKEIFONVDDTLAFWSKH 86

Search completed: May 13, 2003, 14:07:23  
Job time : 16 secs

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21	64	32.2	208	9	US-09-759-1308-355	Sequence 85, Appl
22	64	32.2	208	9	US-10-189-123-85	Sequence 85, Appl
23	64	32.2	225	9	US-09-759-1308-353	Sequence 33, Appl
24	64	32.2	225	9	US-10-189-123-83	Sequence 83, Appl
25	49	24.6	1363	9	US-09-375-248-19	Sequence 19, Appl
26	48	24.1	348	10	US-09-982-610-18	Sequence 18, Appl
27	48	24.1	1298	10	US-09-982-610-33	Sequence 33, Appl
28	48	24.1	1363	9	US-09-375-248-2	Sequence 2, Appl
29	47	23.6	180	10	US-09-766-678-6	Sequence 6, Appl
30	47	23.6	317	9	US-09-939-833-5	Sequence 5, Appl
31	47	23.6	317	10	US-09-939-754-5	Sequence 5, Appl
32	47	23.6	317	10	US-09-939-832-5	Sequence 5, Appl
33	47	23.6	367	9	US-09-939-833-12	Sequence 12, Appl
34	47	23.6	367	10	US-09-939-754-12	Sequence 12, Appl
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36	47	23.6	1356	9	US-09-969-037-7	Sequence 7, Appl
37	47	23.6	1356	9	US-10-022-939-2	Sequence 2, Appl
38	47	23.6	1356	9	US-10-100-405A-2	Sequence 2, Appl
39	47	23.6	1367	10	US-09-919-408-6	Sequence 6, Appl
40	47	23.6	1367	10	US-09-766-678-2	Sequence 2, Appl
41	47	23.6	1367	10	US-09-872-136-6	Sequence 6, Appl
42	44.5	22.4	192	10	US-09-801-368-192	Sequence 192, Appl
43	44.5	22.1	53	10	US-09-982-610-6	Sequence 6, Appl
44	43.5	21.9	49	9	US-09-836-382-34	Sequence 34, Appl
45	43	21.6	100	10	US-09-872-523-9	Sequence 9, Appl

APPLICANT: Grinnell, Brian W

1 TITLE OF INVENTION: PROTEIN C DERIVATIVES

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; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-4

Query Match
Best Local Similarity 91.0%; Score 181; DB 9; Length 419;
Matches 34; Conservative 1; Mismatches 9; Indels 0; Gaps 0;

Db 1 ANSFLLXLRHGSLSRXCIXXICDFXXAKXIFZDVDDTLAFWSKH 44
1 ANSFLEELRHGSLERCEIEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 3
US-10-182-263-5
; Sequence 5, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 5
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-5

Query Match
Best Local Similarity 91.0%; Score 181; DB 9; Length 419;
Matches 34; Conservative 1; Mismatches 9; Indels 0; Gaps 0;

Db 1 ANSFLLXLRHGSLSRXCIXXICDFXXAKXIFZDVDDTLAFWSKH 44
1 ANSFLEELRHGSLERCEIEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 4
US-10-182-263-6
; Sequence 6, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
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; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 6
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-6

Query Match
Best Local Similarity 86.9%; Score 173; DB 9; Length 419;
Matches 33; Conservative 1; Mismatches 10; Indels 0; Gaps 0;

Db 1 ANSFLLXLRHGSLSRXCIXXICDFXXAKXIFZDVDDTLAFWSKH 44
1 ANSFLEELRHGSLERCEIEICDFEAKEIFEDVDDTLAFWSKH 44

RESULT 5
US-10-182-263-1
; Sequence 1, Application US/10182263
; Publication No. US20030022354A1
; GENERAL INFORMATION:
; APPLICANT: Gerlitz, Bruce E
; APPLICANT: Jones, Bryan E
; APPLICANT: Grinnell, Brian W
; TITLE OF INVENTION: PROTEIN C DERIVATIVES
; FILE REFERENCE: X-13611
; CURRENT APPLICATION NUMBER: US/10/182,263
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: 60/181948
; PRIOR FILING DATE: 2002-02-11
; PRIOR APPLICATION NUMBER: 60/189199
; PRIOR FILING DATE: 2000-03-14
; NUMBER OF SEQ ID NOS: 12
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 1
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-182-263-1

Query Match
Best Local Similarity 84.9%; Score 169; DB 9; Length 419;
Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;

Db 1 ANSFLLXLRHGSLSRXCIXXICDFXXAKXIFZDVDDTLAFWSKH 44
1 ANSFLEELRHSSLERCEIEICDFEAKEIFONVDDTLAFWSKH 44

RESULT 6
US-09-978-917A-4
; Sequence 4, Application US/09978917A
; Publication No. US20030027299A1
; GENERAL INFORMATION:
; APPLICANT: Maxygen Aps; Maxygen Holdings
; TITLE OF INVENTION: Protein C or activated protein C-like molecules
; FILE REFERENCE: 0219us310 - protein C
; CURRENT APPLICATION NUMBER: US/09/978,917A
; CURRENT FILING DATE: 2001-10-17
; NUMBER OF SEQ ID NOS: 48
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 4
; LENGTH: 419
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-978-917A-4

Query Match
Best Local Similarity 84.9%; Score 169; DB 9; Length 419;
Matches 32; Conservative 2; Mismatches 10; Indels 0; Gaps 0;

Db 1 ANSFLLXLRHGSLSRXCIXXICDFXXAKXIFZDVDDTLAFWSKH 44
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; PRIOR APPLICATION NUMBER: US 09/420,707
; PRIOR FILING DATE: 1999-10-19
; NUMBER OF SEQ ID NOS: 460
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 313
; LENGTH: 96
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-759-130B-313

Query Match
Best Local Similarity 41.2%; Score 84; DB 9; Length 96;
Matches 14; Conservative 8; Mismatches 12; Indels 0; Gaps 0;

OY 11 GSLRXXCIXXICDFXXKXIFZDVDDTLAFWSKH 44
DB 46 GNLRECNELCNVEARELFVDEDKTIAFWQY 79

RESULT 11
US-10-189-123-43
; Sequence 43, Application US/10189123
; Publication No. US2003008286A1
; GENERAL INFORMATION:
; APPLICANT: KIRST, Susan J.
; APPLICANT: HOLTZMAN, Douglas A.
; APPLICANT: FRASER, Christopher C.
; APPLICANT: SHARP, John D.
; APPLICANT: BARNES, Thomas S.
; TITLE OF INVENTION: ANTIBODIES HAVING DIAGNOSTIC, PREVENTIVE, THERAPEUTIC, AND OTHER
; FILE REFERENCE: 10147-1103
; CURRENT APPLICATION NUMBER: US/10/189,123
; PRIOR FILING DATE: 2002-07-02
; PRIOR APPLICATION NUMBER: US 09/596,194
; PRIOR FILING DATE: 2000-06-16
; PRIOR APPLICATION NUMBER: US 09/342,364
; PRIOR FILING DATE: 1999-06-29
; NUMBER OF SEQ ID NOS: 100
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 43
; LENGTH: 96
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-189-123-43

Query Match
Best Local Similarity 42.2%; Score 84; DB 9; Length 96;
Matches 14; Conservative 8; Mismatches 12; Indels 0; Gaps 0;

OY 11 GSLRXXCIXXICDFXXKXIFZDVDDTLAFWSKH 44
DB 46 GNLRECNELCNVEARELFVDEDKTIAFWQY 79

RESULT 12
US-09-759-130B-312
; Sequence 312, Application US/09759130B
; Publication No. US20030022279A1
; GENERAL INFORMATION:
; APPLICANT: Millennium Pharmaceuticals, Inc.
; APPLICANT: McCarthy, Sean A.
; APPLICANT: Fraser, Christopher C.
; APPLICANT: Sharp, John D.
; APPLICANT: Barnes, Thomas S.
; APPLICANT: Kirst, Susan J.
; APPLICANT: Mackay, Charles R.
; APPLICANT: Myers, Paul S.
; APPLICANT: Leiby, Kevin R.
; APPLICANT: Wrighton, Nicolas
; APPLICANT: Goodheart, Andrew
; APPLICANT: Holtzman, Douglas A.
; TITLE OF INVENTION: NOVEL GENES ENCODING PROTEINS HAVING
; PROGNOSTIC, DIAGNOSTIC, PREVENTIVE, THERAPEUTIC, AND OTHER
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; TITLE OF INVENTION: USES.
; FILE REFERENCE: MP100-535OMNIM
; CURRENT APPLICATION NUMBER: US/09/759,130B
; CURRENT FILING DATE: 2002-09-16
; PRIOR APPLICATION NUMBER: US 09/479,249
; PRIOR FILING DATE: 2000-01-07
; PRIOR APPLICATION NUMBER: US 09/559,497
; PRIOR FILING DATE: 2000-04-27
; PRIOR APPLICATION NUMBER: US 09/578,063
; PRIOR FILING DATE: 2000-05-24
; PRIOR APPLICATION NUMBER: US 09/333,159
; PRIOR FILING DATE: 1999-06-14
; PRIOR APPLICATION NUMBER: US 09/596,194
; PRIOR FILING DATE: 2000-07-14
; PRIOR APPLICATION NUMBER: US 09/342,364
; PRIOR FILING DATE: 1999-06-29
; PRIOR APPLICATION NUMBER: US 09/608,452
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/393,996
; PRIOR FILING DATE: 1999-09-10
; PRIOR APPLICATION NUMBER: US 09/602,871
; PRIOR FILING DATE: 2000-06-23
; PRIOR APPLICATION NUMBER: US 09/420,707
; PRIOR FILING DATE: 1999-10-19
; NUMBER OF SEQ ID NOS: 460
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 312
; LENGTH: 209
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-759-130B-312

Query Match
Best Local Similarity 42.2%; Score 84; DB 9; Length 209;
Matches 14; Conservative 8; Mismatches 12; Indels 0; Gaps 0;

OY 11 GSLRXXCIXXICDFXXKXIFZDVDDTLAFWSKH 44
DB 46 GNLRECNELCNVEARELFVDEDKTIAFWQY 79

RESULT 13
US-10-189-123-42
; Sequence 42, Application US/10189123
; Publication No. US2003008286A1
; GENERAL INFORMATION:
; APPLICANT: KIRST, Susan J.
; APPLICANT: HOLTZMAN, Douglas A.
; APPLICANT: FRASER, Christopher C.
; APPLICANT: SHARP, John D.
; APPLICANT: BARNES, Thomas S.
; TITLE OF INVENTION: ANTIBODIES HAVING DIAGNOSTIC, PREVENTIVE, THERAPEUTIC, AND OTHER
; FILE REFERENCE: 10147-1103
; CURRENT APPLICATION NUMBER: US/10/189,123
; PRIOR FILING DATE: 2002-07-02
; PRIOR APPLICATION NUMBER: US 09/596,194
; PRIOR FILING DATE: 2000-06-16
; PRIOR APPLICATION NUMBER: US 09/342,364
; PRIOR FILING DATE: 1999-06-29
; NUMBER OF SEQ ID NOS: 100
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO: 42
; LENGTH: 209
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-189-123-42

Query Match
Best Local Similarity 42.2%; Score 84; DB 9; Length 209;
Matches 14; Conservative 8; Mismatches 12; Indels 0; Gaps 0;

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